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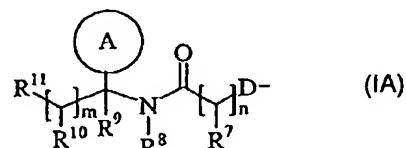
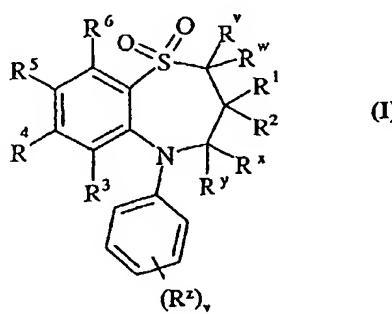
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(54) Title: CHEMICAL COMPOUNDS



(57) Abstract: The present invention relates to compounds of the formula (I): (wherein variable groups are as defined within) pharmaceutically acceptable salts, solvates, solvates of such salts and prodrugs thereof and their use as ileal bile acid transport (IBAT) inhibitors for the treatment of hyperlipidaemia. Processes for their manufacture and pharmaceutical compositions containing them are also described.

WO 02/50051 A1

1,5 BENZOTIAZEPINES AND THEIR USE AS ANTIHYPOLIPIDEMICS

This invention relates to benzothiazepine derivatives, or pharmaceutically acceptable salts, solvates, solvates of such salts and prodrugs thereof. These benzothiazepines possess
5 ileal bile acid transport (IBAT) inhibitory activity and accordingly have value in the treatment of disease states associated with hyperlipidaemic conditions and they are useful in methods of treatment of a warm-blooded animal, such as man. The invention also relates to processes for the manufacture of said benzothiazepine derivatives, to pharmaceutical compositions containing them and to their use in the manufacture of medicaments to inhibit IBAT in a
10 warm-blooded animal, such as man.

It is well-known that hyperlipidaemic conditions associated with elevated concentrations of total cholesterol and low-density lipoprotein cholesterol are major risk factors for cardiovascular atherosclerotic disease (for instance "Coronary Heart Disease: Reducing the Risk; a Worldwide View" Assman G., Carmena R. Cullen P. *et al*; Circulation
15 1999, 100, 1930-1938 and 'Diabetes and Cardiovascular Disease: A Statement for Healthcare Professionals from the American Heart Association' Grundy S, Benjamin I., Burke G., *et al*; Circulation, 1999, 100, 1134-46). Interfering with the circulation of bile acids within the lumen of the intestinal tracts is found to reduce the level of cholesterol. Previous established therapies to reduce the concentration of cholesterol involve, for instance, treatment with
20 HMG-CoA reductase inhibitors, preferably statins such as simvastatin and fluvastatin, or treatment with bile acid binders, such as resins. Frequently used bile acid binders are for instance cholestyramine and colestipol. One recently proposed therapy ("Bile Acids and Lipoprotein Metabolism: a Renaissance for Bile Acids in the Post Statin Era" Angelin B, Eriksson M, Rudling M; Current Opinion on Lipidology, 1999, 10, 269-74) involved the
25 treatment with substances with an IBAT inhibitory effect.

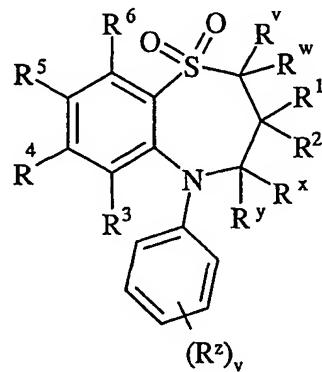
Re-absorption of bile acid from the gastro-intestinal tract is a normal physiological process which mainly takes place in the ileum by the IBAT mechanism. Inhibitors of IBAT can be used in the treatment of hypercholesterolaemia (see for instance "Interaction of bile acids and cholesterol with nonsystemic agents having hypcholesterolaemic properties",
30 Biochimica et Biophysica Acta, 1210 (1994) 255- 287). Thus, suitable compounds having such inhibitory IBAT activity are also useful in the treatment of hyperlipidaemic conditions. Compounds possessing such IBAT inhibitory activity have been described, see for instance compounds described in WO 93/16055, WO 94/18183, WO 94/18184, WO 96/05188, WO

96/08484, WO 96/16051, WO 97/33882, WO 98/38182, WO 99/35135, WO 98/40375, WO 99/35153, WO 99/64409, WO 99/64410, WO 00/01687, WO 00/47568, WO 00/61568, WO 01/68906, DE 19825804, WO 00/38725, WO 00/38726, WO 00/38727, WO 00/38728, WO 00/38729, WO 01/68906, and EP 0 864 582.

5 A further aspect of this invention relates to the use of the compounds of the invention in the treatment of dyslipidemic conditions and disorders such as hyperlipidaemia, hypertriglyceridemia, hyperbetalipoproteinemia (high LDL), hyperprebetalipoproteinemia (high VLDL), hyperchylomicronemia, hypolipoproteinemia, hypercholesterolemia, hyperlipoproteinemia and hypoalphalipoproteinemia (low HDL). In addition, these
10 compounds are expected to be useful for the prevention and treatment of different clinical conditions such as atherosclerosis, arteriosclerosis, arrhythmia, hyper-thrombotic conditions, vascular dysfunction, endothelial dysfunction, heart failure, coronary heart diseases, cardiovascular diseases, myocardial infarction, angina pectoris, peripheral vascular diseases, inflammation of cardiovascular tissues such as heart, valves, vasculature, arteries and veins,
15 aneurisms, stenosis, restenosis, vascular plaques, vascular fatty streaks, leukocyte, monocytes and/or macrophage infiltrate, intimal thickening, medial thinning, infectious and surgical trauma and vascular thrombosis, stroke and transient ischaemic attacks.

The present invention is based on the discovery that certain benzothiazepine compounds surprisingly inhibit IBAT. Such properties are expected to be of value in the
20 treatment of disease states associated with hyperlipidaemic conditions.

Accordingly, the present invention provides a compound of formula (I):



(I)

wherein:

25 R^v and R^w are independently selected from hydrogen or C₁₋₆alkyl;
 R^1 and R^2 are independently selected from C₁₋₆alkyl;

R^x and R^y are independently selected from hydrogen or C₁₋₆alkyl, or one of R^x and R^y is hydrogen or C₁₋₆alkyl and the other is hydroxy or C₁₋₆alkoxy;

R^z is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkanoyl, C₁₋₆alkanoyloxy,

5 N -(C₁₋₆alkyl)amino, N,N -(C₁₋₆alkyl)₂amino, C₁₋₆alkanoylamino, N -(C₁₋₆alkyl)carbamoyl,

N,N -(C₁₋₆alkyl)₂carbamoyl, C₁₋₆alkylS(O)_a wherein a is 0 to 2, C₁₋₆alkoxycarbonyl,

C₁₋₆alkoxycarbonylamino, ureido, N' -(C₁₋₆alkyl)ureido, N -(C₁₋₆alkyl)ureido,

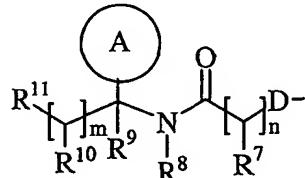
N',N' -(C₁₋₆alkyl)₂ureido, N' -(C₁₋₆alkyl)- N -(C₁₋₆alkyl)ureido,

N',N' -(C₁₋₆alkyl)₂- N -(C₁₋₆alkyl)ureido, N -(C₁₋₆alkyl)sulphamoyl and

10 N,N -(C₁₋₆alkyl)₂sulphamoyl;

v is 0-5;

one of R^4 and R^5 is a group of formula (IA):



(IA)

15 R^3 and R^6 and the other of R^4 and R^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N -(C₁₋₄alkyl)amino, N,N -(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N -(C₁₋₄alkyl)carbamoyl, N,N -(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein a is 0 to 2, C₁₋₄alkoxycarbonyl,

20 N -(C₁₋₄alkyl)sulphamoyl and N,N -(C₁₋₄alkyl)₂sulphamoyl; wherein R^3 and R^6 and the other of R^4 and R^5 may be optionally substituted on carbon by one or more R^{16} ;

D is -O-, -N(R^a)-, -S(O)_b- or -CH(R^a)-; wherein R^a is hydrogen or C₁₋₆alkyl and b is

0-2;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more
25 substituents selected from R^{17} ;

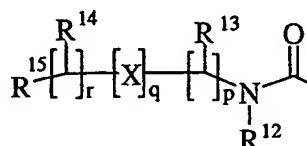
R^7 is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R^7 is optionally substituted by one or more substituents selected from R^{18} ;

R^8 is hydrogen or C₁₋₄alkyl;

R^9 is hydrogen or C₁₋₄alkyl;

R^{10} is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R^{10} is optionally substituted by one or more substituents selected from R^{19} ;

R^{11} is carboxy, sulpho, sulphino, phosphono, tetrazolyl, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently selected from C₁₋₆alkyl; or R¹¹ is a group of formula (IB):



(IB)

wherein:

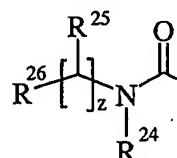
X is -N(R^q)-, -N(R^q)C(O)-, -O-, and -S(O)_a-; wherein a is 0-2 and R^q is hydrogen or C₁₋₄alkyl;

R^{12} is hydrogen or C₁₋₄alkyl;

R^{13} and R^{14} are independently selected from hydrogen, C₁₋₄alkyl, carbocyclyl, heterocyclyl or R²³; wherein said C₁₋₄alkyl, carbocyclyl or heterocyclyl may be independently optionally substituted by one or more substituents selected from R²⁰;

R^{15} is carboxy, sulpho, sulphino, phosphono, tetrazolyl, -P(O)(OR^e)(OR^f),

-P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₆alkyl; or R¹⁵ is a group of formula (IC):



(IC)

20 wherein:

R^{24} is selected from hydrogen or C₁₋₄alkyl;

R^{25} is selected from hydrogen, C₁₋₄alkyl, carbocyclyl, heterocyclyl or R²⁷; wherein said C₁₋₄alkyl, carbocyclyl or heterocyclyl may be independently optionally substituted by one or more substituents selected from R²⁸;

R^{26} is selected from carboxy, sulpho, sulphino, phosphono, tetrazolyl,

-P(O)(OR^g)(OR^h), -P(O)(OH)(OR^g), -P(O)(OH)(R^g) or -P(O)(OR^g)(R^h) wherein R^g and R^h are independently selected from C₁₋₆alkyl;

p is 1-3; wherein the values of R¹³ may be the same or different;

q is 0-1;

r is 0-3; wherein the values of R¹⁴ may be the same or different;

m is 0-2; wherein the values of R¹⁰ may be the same or different;

n is 1-3; wherein the values of R⁷ may be the same or different;

5 z is 0-3; wherein the values of R²⁵ may be the same or different;

R¹⁶, R¹⁷ and R¹⁸ are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a

10 wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and

N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R¹⁶, R¹⁷ and R¹⁸ may be independently optionally substituted on carbon by one or more R²¹;

R¹⁹, R²⁰, R²³, R²⁷ and R²⁸ are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl,

15 C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino,

C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl,

N,N-(C₁₋₄alkyl)₂sulphamoyl, carbocyclyl, heterocyclyl, sulpho, sulphino, amidino, phosphono, -P(O)(OR^a)(OR^b), -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or -P(O)(OR^a)(R^b), wherein R^a and R^b are

20 independently selected from C₁₋₆alkyl; wherein R¹⁹, R²⁰, R²³, R²⁷ and R²⁸ may be independently optionally substituted on carbon by one or more R²²;

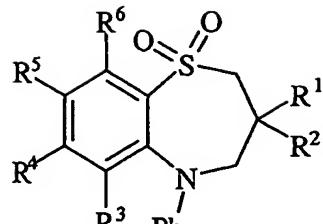
R²¹ and R²² are independently selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl,

25 formamido, acetylarnino, acetoxy, methylarnino, dimethylarnino, N-methylcarbamoyl, N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl and N,N-dimethylsulphamoyl;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

According to a further aspect of the present invention there is provided a compound of

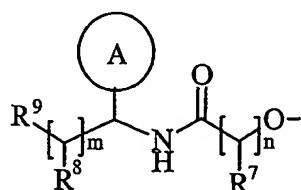
30 formula (I'):



(I')

wherein:

5 \mathbf{R}^1 and \mathbf{R}^2 are independently selected from C₁₋₆alkyl;
 one of \mathbf{R}^4 and \mathbf{R}^5 is a group of formula (IA'):

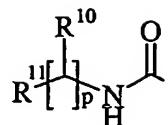


(IA')

\mathbf{R}^3 and \mathbf{R}^6 and the other of \mathbf{R}^4 and \mathbf{R}^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl,
 10 C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R³ and R⁶ and the other of R⁴ and R⁵ may be optionally substituted on carbon by one or more R¹²;

15 Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R¹³;
 R⁷ is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R⁷ is optionally substituted by one or more substituents selected from R¹⁴;

R⁸ is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R⁸ is optionally substituted by one or more substituents selected from R¹⁵;
 20 R⁹ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently selected from C₁₋₆alkyl; or R⁹ is a group of formula (IB'):



(IB')

wherein:

R¹⁰ is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R¹⁰ is optionally
5 substituted by one or more substituents selected from R¹⁶;

R¹¹ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e),
-P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from
C₁₋₆alkyl;

- p** is 1-3; wherein the values of R¹⁰ may be the same or different;
- 10 **m** is 0-2; wherein the values of R⁸ may be the same or different;
- n** is 1-3; wherein the values of R⁷ may be the same or different;
- R¹²**, **R¹³** and **R¹⁴** are independently selected from halo, nitro, cyano, hydroxy, amino,
carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy,
C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, *N*-(C₁₋₄alkyl)amino, *N,N*-(C₁₋₄alkyl)₂amino,
15 C₁₋₄alkanoylamino, *N*-(C₁₋₄alkyl)carbamoyl, *N,N*-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a
wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, *N*-(C₁₋₄alkyl)sulphamoyl and
N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R¹², R¹³ and R¹⁴ may be independently optionally
substituted on carbon by one or more R¹⁷;

- R¹⁵** and **R¹⁶** are independently selected from halo, nitro, cyano, hydroxy, amino,
20 carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy,
C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, *N*-(C₁₋₄alkyl)amino, *N,N*-(C₁₋₄alkyl)₂amino,
C₁₋₄alkanoylamino, *N*-(C₁₋₄alkyl)carbamoyl, *N,N*-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a
wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, *N*-(C₁₋₄alkyl)sulphamoyl and
N,N-(C₁₋₄alkyl)₂sulphamoyl, sulpho, sulphino, amidino, phosphono, -P(O)(OR^a)(OR^b),

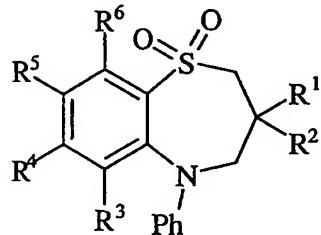
- 25 -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or -P(O)(OR^a)(R^b), wherein R^a and R^b are independently
selected from C₁₋₆alkyl; wherein R¹⁵ and R¹⁶ may be independently optionally substituted on
carbon by one or more R¹⁸;

- R¹⁷** and **R¹⁸** are independently selected from halo, hydroxy, cyano, carbamoyl, ureido,
amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy,
30 methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl,
formamido, acetylamino, acetoxy, methylamino, dimethylamino, *N*-methylcarbamoyl,

N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, *N*-methylsulphamoyl and *N,N*-dimethylsulphamoyl;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

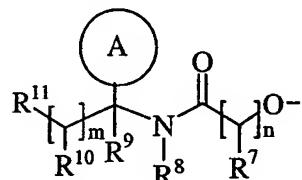
According to a further aspect of the present invention there is provided a compound of
5 formula (I''):



(I'')

wherein:

10 \mathbf{R}^1 and \mathbf{R}^2 are independently selected from C₁₋₆alkyl;
one of \mathbf{R}^4 and \mathbf{R}^5 is a group of formula (IA''):



(IA'')

15 \mathbf{R}^3 and \mathbf{R}^6 and the other of \mathbf{R}^4 and \mathbf{R}^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, *N*-(C₁₋₄alkyl)amino, *N,N*-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, *N*-(C₁₋₄alkyl)carbamoyl, *N,N*-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and *N,N*-(C₁₋₄alkyl)₂sulphamoyl; wherein \mathbf{R}^3 and \mathbf{R}^6 and the other of \mathbf{R}^4 and \mathbf{R}^5 may be optionally substituted on carbon by one or more \mathbf{R}^{16} ;

20 Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from \mathbf{R}^{17} ;

\mathbf{R}^7 is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein \mathbf{R}^7 is optionally substituted by one or more substituents selected from \mathbf{R}^{18} ;

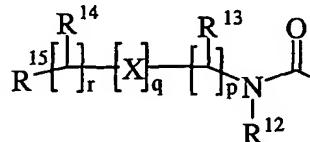
\mathbf{R}^8 is hydrogen or C₁₋₄alkyl;

25 \mathbf{R}^9 is hydrogen or C₁₋₄alkyl;

R¹⁰ is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R¹⁰ is optionally substituted by one or more substituents selected from R¹⁹;

R¹¹ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently selected from

5 C₁₋₆alkyl; or R¹¹ is a group of formula (IB''):



(IB'')

wherein:

X is -N(R^q)-, -N(R^q)C(O)-, -O-, and -S(O)_a-; wherein a is 0-2 and R^q is hydrogen or
10 C₁₋₄alkyl;

R¹² is hydrogen or C₁₋₄alkyl;

R¹³ and R¹⁴ are independently selected from hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R¹³ and R¹⁴ may be independently optionally substituted by one or more substituents selected from R²⁰;

15 R¹⁵ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₆alkyl;

p is 1-3; wherein the values of R¹³ may be the same or different;

q is 0-1;

20 r is 0-3; wherein the values of R¹⁴ may be the same or different;

m is 0-2; wherein the values of R¹⁰ may be the same or different;

n is 1-3; wherein the values of R⁷ may be the same or different;

R¹⁶, R¹⁷ and R¹⁸ are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy,

25 C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino,

C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a

wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and

N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R¹⁶, R¹⁷ and R¹⁸ may be independently optionally substituted on carbon by one or more R²¹;

30 R¹⁹ and R²⁰ are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy,

C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino,
C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a
wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl,
N,N-(C₁₋₄alkyl)₂sulphamoyl, carbocyclyl, heterocyclyl, sulpho, sulphino, amidino, phosphono,
5 -P(O)(OR^a)(OR^b), -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or -P(O)(OR^a)(R^b), wherein R^a and R^b are
independently selected from C₁₋₆alkyl; wherein R¹⁹ and R²⁰ may be independently optionally
substituted on carbon by one or more R²²;

R²¹ and R²² are independently selected from halo, hydroxy, cyano, carbamoyl, ureido,
amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy,
10 methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl,
formamido, acetylarnino, acetoxy, methylarnino, dimethylarnino, N-methylcarbamoyl,
N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl and
N,N-dimethylsulphamoyl;
or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

15 In the following paragraphs of the description, and in the claims, where a compound
of formula (I) is referred to, it is to be understood that this aspect also relates to compounds of
formula (I') and compounds of formula (I'').

In addition, the skilled person will appreciate that the numbering system differs
between compounds of formula (I) and compounds of formula (I'). The numbering system
20 used hereinbelow refers to compounds of formula (I), but it is to be understood that these
statements also apply to the corresponding values in formula (I').

In this specification the term "alkyl" includes both straight and branched chain alkyl
groups but references to individual alkyl groups such as "propyl" are specific for the straight
chain version only. For example, "C₁₋₆alkyl" includes C₁₋₄alkyl, C₁₋₃alkyl, propyl, isopropyl
25 and t-butyl. However, references to individual alkyl groups such as 'propyl' are specific for
the straight chained version only and references to individual branched chain alkyl groups
such as 'isopropyl' are specific for the branched chain version only. A similar convention
applies to other radicals, for example "phenylC₁₋₆alkyl" would include phenylC₁₋₄alkyl,
benzyl, 1-phenylethyl and 2-phenylethyl. The term "halo" refers to fluoro, chloro, bromo and
30 iodo.

Where optional substituents are chosen from "one or more" groups it is to be
understood that this definition includes all substituents being chosen from one of the specified
groups or the substituents being chosen from two or more of the specified groups.

"Heteroaryl" is a totally unsaturated, mono or bicyclic ring containing 3-12 atoms of which at least one atom is chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked. Preferably "heteroaryl" refers to a totally unsaturated, monocyclic ring containing 5 or 6 atoms or a bicyclic ring containing 9 or 10 atoms of which at least one atom is chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked. In another aspect of the invention, "heteroaryl" refers to a totally unsaturated, monocyclic ring containing 5 or 6 atoms or a bicyclic ring containing 8, 9 or 10 atoms of which at least one atom is chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked.

5 Examples and suitable values of the term "heteroaryl" are thienyl, isoxazolyl, imidazolyl, pyrrolyl, thiadiazolyl, isothiazolyl, triazolyl, pyranyl, indolyl, pyrimidyl, pyrazinyl, pyridazinyl, pyridyl and quinolyl. Preferably the term "heteroaryl" refers to thienyl or indolyl.

"Aryl" is a totally unsaturated, mono or bicyclic carbon ring that contains 3-12 atoms. Preferably "aryl" is a monocyclic ring containing 5 or 6 atoms or a bicyclic ring containing 9 or 10 atoms. Suitable values for "aryl" include phenyl or naphthyl. Particularly "aryl" is phenyl.

A "heterocyclyl" is a saturated, partially saturated or unsaturated, mono or bicyclic ring containing 3-12 atoms of which at least one atom is chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked, wherein a -CH₂- group can optionally be replaced by a -C(O)- or a ring sulphur atom may be optionally oxidised to form the S-oxides. Preferably a "heterocyclyl" is a saturated, partially saturated or unsaturated, mono or bicyclic ring containing 5 or 6 atoms of which at least one atom is chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked, wherein a -CH₂- group can optionally be replaced by a -C(O)- or a ring sulphur atom may be optionally oxidised to form S-oxide(s). Examples and suitable values of the term "heterocyclyl" are thiazolidinyl, pyrrolidinyl, pyrrolinyl, 2-pyrrolidonyl, 2,5-dioxopyrrolidinyl, 2-benzoxazolinonyl, 1,1-dioxotetrahydrothienyl, 2,4-dioxoimidazolidinyl, 2-oxo-1,3,4-(4-triazolinyl), 2-oxazolidinonyl, 5,6-dihydouracilyl, 1,3-benzodioxolyl, 1,2,4-oxadiazolyl, 2-azabicyclo[2.2.1]heptyl, 4-thiazolidonyl, morpholino, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, 2,3-dihydrobenzofuranyl, benzothienyl, tetrahydropyrananyl, piperidyl, 1-oxo-1,3-dihydroisoindolyl, piperazinyl, thiomorpholino, 1,1-dioxothiomorpholino, tetrahydropyrananyl, 1,3-dioxolanyl, homopiperazinyl, thienyl, isoxazolyl, imidazolyl, pyrrolyl, thiadiazolyl, isothiazolyl, 1,2,4-triazolyl, 1,3,4-triazolyl,

pyranyl, indolyl, pyrimidyl, thiazolyl, pyrazinyl, pyridazinyl, pyridyl, 4-pyridonyl, quinolyl and 1-isoquinolonyl.

A “carbocyclyl” is a saturated, partially saturated or unsaturated, mono or bicyclic carbon ring that contains 3-12 atoms; wherein a -CH₂- group can optionally be replaced by a 5 -C(O)-. Preferably “carbocyclyl” is a monocyclic ring containing 5 or 6 atoms or a bicyclic ring containing 9 or 10 atoms. Suitable values for “carbocyclyl” include cyclopropyl, cyclobutyl, 1-oxocyclopentyl, cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl, phenyl, naphthyl, tetralinyl, indanyl or 1-oxoindanyl. Particularly “carbocyclyl” is cyclopropyl, cyclobutyl, 1-oxocyclopentyl, cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl, phenyl 10 or 1-oxoindanyl.

An example of “C₁₋₆alkanoyloxy” and “C₁₋₄alkanoyloxy” is acetoxy. Examples of “C₁₋₆alkoxycarbonyl” and “C₁₋₄alkoxycarbonyl” include methoxycarbonyl, ethoxycarbonyl, *n*- and *t*-butoxycarbonyl. Examples of “C₁₋₆alkoxy” and “C₁₋₄alkoxy” include methoxy, ethoxy and propoxy. Examples of “C₁₋₆alkanoylamino” and “C₁₋₄alkanoylamino” include formamido, 15 acetamido and propionylamino. Examples of “C₁₋₆alkylS(O)_a wherein a is 0 to 2” and “C₁₋₄alkylS(O)_a wherein a is 0 to 2” include methylthio, ethylthio, methylsulphinyl, ethylsulphinyl, mesyl and ethylsulphonyl. Examples of “C₁₋₆alkanoyl” and “C₁₋₄alkanoyl” include C₁₋₃alkanoyl, propionyl and acetyl. Examples of “N-(C₁₋₆alkyl)amino” and “N-(C₁₋₄alkyl)amino” include methylamino and ethylamino. Examples of 20 “N,N-(C₁₋₆alkyl)₂amino” and “N,N-(C₁₋₄alkyl)₂amino” include di-N-methylamino, di-(*N*-ethyl)amino and *N*-ethyl-*N*-methylamino. Examples of “C₂₋₆alkenyl” and “C₂₋₄alkenyl” are vinyl, allyl and 1-propenyl. Examples of “C₂₋₆alkynyl” and “C₂₋₄alkynyl” are ethynyl, 1-propynyl and 2-propynyl. Examples of “N-(C₁₋₆alkyl)sulphamoyl” and “N-(C₁₋₄alkyl)sulphamoyl” are *N*-(C₁₋₃alkyl)sulphamoyl, *N*-(methyl)sulphamoyl and 25 N-(ethyl)sulphamoyl. Examples of “N-(C₁₋₆alkyl)₂sulphamoyl” and “N-(C₁₋₄alkyl)₂sulphamoyl” are *N,N*-(dimethyl)sulphamoyl and *N*-(methyl)-*N*-(ethyl)sulphamoyl. Examples of “N-(C₁₋₆alkyl)carbamoyl” and “N-(C₁₋₄alkyl)carbamoyl” are methylaminocarbonyl and ethylaminocarbonyl. Examples of “N,N-(C₁₋₆alkyl)₂carbamoyl” and “N,N-(C₁₋₄alkyl)₂carbamoyl” are dimethylaminocarbonyl 30 and methylethylaminocarbonyl. Examples of “C₁₋₆alkoxycarbonylamino” are ethoxycarbonylamino and *t*-butoxycarbonylamino. Examples of “N’-(C₁₋₆alkyl)ureido” are *N*'-methylureido and *N*'-ethylureido. Examples of “N-(C₁₋₆alkyl)ureido” are *N*-methylureido and *N*-ethylureido. Examples of “N’,N’-(C₁₋₆alkyl)₂ureido” are *N’,N'*-dimethylureido and *N'*-

methyl-*N'*-ethylureido. Examples of "*N'*-(C₁₋₆alkyl)-*N*-(C₁₋₆alkyl)ureido are *N'*-methyl-*N*-methylureido and *N'*-propyl-*N*-methylureido. Examples of "*N',N'*-(C₁₋₆alkyl)₂-*N*-(C₁₋₆alkyl)ureido are *N',N'*-dimethyl-*N*-methylureido and *N'*-methyl-*N'*-ethyl-*N*-propylureido.

5 A suitable pharmaceutically acceptable salt of a compound of the invention is, for example, an acid-addition salt of a compound of the invention which is sufficiently basic, for example, an acid-addition salt with, for example, an inorganic or organic acid, for example hydrochloric, hydrobromic, sulphuric, phosphoric, trifluoroacetic, citric or maleic acid. In addition a suitable pharmaceutically acceptable salt of a compound of the invention which is 10 sufficiently acidic is an alkali metal salt, for example a sodium or potassium salt, an alkaline earth metal salt, for example a calcium or magnesium salt, an ammonium salt or a salt with an organic base which affords a physiologically-acceptable cation, for example a salt with methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.

15 The compounds of the formula (I) may be administered in the form of a pro-drug which is broken down in the human or animal body to give a compound of the formula (I). Examples of pro-drugs include *in vivo* hydrolysable esters and *in vivo* hydrolysable amides of a compound of the formula (I).

An *in vivo* hydrolysable ester of a compound of the formula (I) containing carboxy or 20 hydroxy group is, for example, a pharmaceutically acceptable ester which is hydrolysed in the human or animal body to produce the parent acid or alcohol. Suitable pharmaceutically acceptable esters for carboxy include C₁₋₆alkoxymethyl esters for example methoxymethyl, C₁₋₆alkanoyloxymethyl esters for example pivaloyloxymethyl, phthalidyl esters, C₃₋₈cycloalkoxycarbonyloxyC₁₋₆alkyl esters for example 1-cyclohexylcarbonyloxyethyl; 25 1,3-dioxolen-2-onylmethyl esters for example 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁₋₆alkoxycarbonyloxyethyl esters for example 1-methoxycarbonyloxyethyl and may be formed at any carboxy group in the compounds of this invention.

An *in vivo* hydrolysable ester of a compound of the formula (I) containing a hydroxy group includes inorganic esters such as phosphate esters and α -acyloxyalkyl ethers and 30 related compounds which as a result of the *in vivo* hydrolysis of the ester breakdown to give the parent hydroxy group. Examples of α -acyloxyalkyl ethers include acetoxymethoxy and 2,2-dimethylpropionyloxy-methoxy. A selection of *in vivo* hydrolysable ester forming groups for hydroxy include alkanoyl, benzoyl, phenylacetyl and substituted benzoyl and

phenylacetyl, alkoxycarbonyl (to give alkyl carbonate esters), dialkylcarbamoyl and *N*-(dialkylaminoethyl)-*N*-alkylcarbamoyl (to give carbamates), dialkylaminoacetyl and carboxyacetyl. Examples of substituents on benzoyl include morpholino and piperazino linked from a ring nitrogen atom via a methylene group to the 3- or 4- position of the benzoyl ring.

5 A suitable value for an *in vivo* hydrolysable amide of a compound of the formula (I) containing a carboxy group is, for example, a *N*-C₁₋₆alkyl or *N,N*-di-C₁₋₆alkyl amide such as *N*-methyl, *N*-ethyl, *N*-propyl, *N,N*-dimethyl, *N*-ethyl-*N*-methyl or *N,N*-diethyl amide.

Some compounds of the formula (I) may have chiral centres and/or geometric isomeric centres (E- and Z-isomers), and it is to be understood that the invention 10 encompasses all such optical, diastereoisomers and geometric isomers that possess IBAT inhibitory activity.

The invention relates to any and all tautomeric forms of the compounds of the formula (I) that possess IBAT inhibitory activity.

It is also to be understood that certain compounds of the formula (I) can exist in 15 solvated as well as unsolvated forms such as, for example, hydrated forms. It is to be understood that the invention encompasses all such solvated forms which possess IBAT inhibitory activity.

Preferred values of R¹, R², R³, R⁴, R⁵ and R⁶ are as follows. Such values may be used where appropriate with any of the definitions, claims or embodiments defined hereinbefore or 20 hereinafter.

Preferably R^y and R^w are both hydrogen.

Preferably R¹ and R² are independently selected from C₁₋₄alkyl.

More preferably R¹ and R² are independently selected from ethyl or butyl.

More preferably R¹ and R² are independently selected from ethyl, propyl or butyl.

25 In one aspect of the invention particularly R¹ and R² are both butyl.

In a further aspect of the invention particularly R¹ and R² are both propyl.

In another aspect of the invention particularly one of R¹ and R² is ethyl and the other is butyl.

Preferably R^x and R^y are independently selected from hydrogen or C₁₋₆alkyl.

30 More preferably R^x and R^y are both hydrogen.

Preferably R^z is selected from halo, amino, C₁₋₆alkyl, C₁₋₆alkoxycarbonylamino or N'-(C₁₋₆alkyl)ureido.

More preferably R² is selected from chloro, amino, *t*-butyl, *t*-butoxycarbonylamino or *N'*-(*t*-butyl)ureido.

Preferably v is 0 or 1.

In one aspect of the invention, more preferably v is 0.

5 In one aspect of the invention, more preferably v is 1.

In one aspect of the invention preferably R⁴ is a group of formula (IA) (as depicted above).

In another aspect of the invention preferably R⁵ is a group of formula (IA) (as depicted above).

10 Preferably R³ and R⁶ are hydrogen.

Preferably the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from halo, C₁₋₄alkoxy or C₁₋₄alkylS(O)_a wherein a is 0 to 2; wherein that R⁴ or R⁵ may be optionally substituted on carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy and *N,N*-(C₁₋₄alkyl)₂amino.

15 More preferably the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from bromo, methoxy, isopropoxy, methylthio, ethylthio, isopropylthio or mesyl; wherein that R⁴ or R⁵ may be optionally substituted on carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy and *N,N*-dimethylamino.

Particularly the other of R⁴ and R⁵ that is not the group of formula (IA) is selected
20 from bromo, methoxy, isopropoxy, methylthio, ethylthio, isopropylthio, 2-hydroxyethylthio,
2-(*N,N*-dimethylamino)ethylthio or mesyl.

More particularly the other of R⁴ and R⁵ that is not the group of formula (IA) is methylthio.

Preferably the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from
25 hydrogen, halo, C₁₋₄alkoxy or C₁₋₄alkylS(O)_a wherein a is 0 to 2; wherein that R⁴ or R⁵ may be optionally substituted on carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy, carboxy and *N,N*-(C₁₋₄alkyl)₂amino.

More preferably the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from hydrogen, bromo, methoxy, isopropoxy, methylthio, ethylthio, isopropylthio or mesyl;
30 wherein that R⁴ or R⁵ may be optionally substituted on carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy, carboxy and *N,N*-dimethylamino.

Particularly the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from hydrogen, bromo, methoxy, isopropoxy, methylthio, carboxymethylthio, ethylthio, isopropylthio, 2-hydroxyethylthio, 2-(N,N-dimethylamino)ethylthio or mesyl.

In another aspect of the invention, more preferably the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from hydrogen, chloro, bromo, methoxy, isopropoxy, methylthio, ethylthio or isopropylthio; wherein that R⁴ or R⁵ may be optionally substituted on carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy, carboxy and N,N-dimethylamino.

In another aspect of the invention, particularly the other of R⁴ and R⁵ that is not the group of formula (IA) is selected from hydrogen, chloro, bromo, methoxy, isopropoxy, methylthio, carboxymethylthio, ethylthio, isopropylthio, 2-hydroxyethylthio or 2-(N,N-dimethylamino)ethylthio.

In another aspect of the invention, more particularly the other of R⁴ and R⁵ that is not the group of formula (IA) is bromo or chloro.

In another aspect of the invention, more particularly the other of R⁴ and R⁵ that is not the group of formula (IA) is methoxy.

In one aspect of the invention, preferably Ring A is aryl.

In another aspect of the invention, preferably Ring A is heteroaryl.

When Ring A is aryl, preferably Ring A is phenyl.

When Ring A is heteroaryl, preferably Ring A is thienyl or indolyl.

Preferably Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R¹⁷; wherein

R¹⁷ is selected from halo, hydroxy or C₁₋₄alkyl; wherein R¹⁷ may be optionally substituted on carbon by one or more R²¹; wherein

R²¹ is selected from halo.

Preferably D is -O- or -S-.

In one aspect of the invention, more preferably D is -O-.

In one aspect of the invention, more preferably D is -S-.

More preferably Ring A is phenyl, thienyl or indolyl; wherein Ring A is optionally substituted by one or more substituents selected from halo, hydroxy or trifluoromethyl.

Particularly Ring A is selected from phenyl, 4-hydroxyphenyl, thien-2-yl, 4-trifluoromethylphenyl, 3-hydroxyphenyl, 2-fluorophenyl, 2,3-dihydroxyphenyl or indol-3-yl.

More particularly Ring A is phenyl.

In another aspect of the invention, preferably Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R¹⁷; wherein

R¹⁷ is selected from halo, hydroxy, C₁₋₄alkyl or C₁₋₄alkoxy; wherein R¹⁷ may be

5 optionally substituted on carbon by one or more R²¹; wherein

R²¹ is selected from halo.

In another aspect of the invention, more preferably Ring A is phenyl, thienyl or indolyl; wherein Ring A is optionally substituted by one or more substituents selected from halo, hydroxy, methoxy or trifluoromethyl.

10 In another aspect of the invention, particularly Ring A is selected from phenyl, 4-hydroxyphenyl, 4-methoxyphenyl, thien-2-yl, 4-trifluoromethylphenyl, 3-hydroxyphenyl, 2-fluorophenyl, 2,3-dihydroxyphenyl or indol-3-yl.

In a further aspect of the invention, particularly Ring A is selected from phenyl, 4-hydroxyphenyl, 4-methoxyphenyl, thien-2-yl, 4-trifluoromethylphenyl, 3-hydroxyphenyl, 15 2-fluorophenyl, 4-fluorophenyl, 2,3-dihydroxyphenyl or indol-3-yl.

Preferably R⁷ is hydrogen, C₁₋₄alkyl or carbocyclyl.

More preferably R⁷ is hydrogen, methyl or phenyl.

Particularly R⁷ is hydrogen.

In one aspect of the invention, preferably R⁸ is hydrogen.

20 In another aspect of the invention, preferably R⁸ is C₁₋₄alkyl.

In another aspect of the invention, more preferably R⁸ is hydrogen or methyl.

In one aspect of the invention, preferably R⁹ is hydrogen.

In another aspect of the invention, preferably R⁹ is C₁₋₄alkyl.

In another aspect of the invention, more preferably R⁹ is hydrogen or methyl.

25 Preferably R¹⁰ is hydrogen.

In one aspect of the invention, preferably R¹¹ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently selected from C₁₋₆alkyl.

30 In another aspect of the invention, preferably R¹¹ is a group of formula (IB) (as depicted above).

Preferably R¹¹ is carboxy, -P(O)(OH)(OR^c) or a group of formula (IB) (as depicted above).

More preferably R¹¹ is carboxy, -P(O)(OH)(OEt) or a group of formula (IB) (as depicted above).

In another aspect of the invention, preferably R¹¹ is carboxy, sulpho, -P(O)(OH)(OR^c) wherein R^c is selected from C₁₋₄alkyl or a group of formula (IB) (as depicted above).

5 Preferably X is -NH- or -NHC(O)-.

More preferably X is -NHC(O)-.

In one aspect of the invention, preferably R¹² is hydrogen.

In another aspect of the invention, preferably R¹² is C₁₋₄alkyl.

In another aspect of the invention, more preferably R¹² is hydrogen or methyl.

10 Preferably R¹³ is hydrogen, C₁₋₄alkyl or carbocyclyl; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein R²⁰ is hydroxy.

More preferably R¹³ is hydrogen, methyl or phenyl; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein

15 R²⁰ is hydroxy.

Particularly R¹³ is hydrogen, hydroxymethyl or phenyl.

More particularly R¹³ is hydrogen or hydroxymethyl.

In another aspect of the invention, preferably R¹³ is hydrogen, C₁₋₄alkyl or carbocyclyl; wherein R¹³ is optionally substituted by one or more substituents selected from 20 R²⁰; wherein

R²⁰ is hydroxy, carboxy, carbocyclyl or amino; wherein R²⁰ may be optionally substituted on carbon by one or more R²²;

R²² is hydroxy.

25 In another aspect of the invention, more preferably R¹³ is hydrogen, methyl, ethyl, butyl or phenyl; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein

R²⁰ is hydroxy, carboxy, phenyl or amino; wherein R²⁰ may be optionally substituted on carbon by one or more R²²;

R²² is hydroxy.

30 In another aspect of the invention, particularly R¹³ is hydrogen, hydroxymethyl, 4-aminobutyl, 2-carboxyethyl, 4-hydroxybenzyl or phenyl.

In a further aspect of the invention, preferably R¹³ is hydrogen, C₁₋₄alkyl or carbocyclyl; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein

R²⁰ is hydroxy, carboxy, carbocyclyl, heterocyclyl or amino; wherein R²⁰ may be
5 optionally substituted on carbon by one or more R²²;
R²² is hydroxy.

In a further aspect of the invention, more preferably R¹³ is hydrogen, methyl, ethyl, butyl or phenyl; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein

10 R²⁰ is hydroxy, carboxy, phenyl, imidazolyl or amino; wherein R²⁰ may be optionally substituted on carbon by one or more R²²;
R²² is hydroxy.

In a further aspect of the invention, particularly R¹³ is hydrogen, hydroxymethyl, 4-aminobutyl, 2-carboxyethyl, 4-hydroxybenzyl, imidazol-5-ylmethyl or phenyl.

15 In another further aspect of the invention, preferably R¹³ is hydrogen, C₁₋₄alkyl, carbocyclyl or R²³; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein

R²⁰ is hydroxy, C₁₋₄alkylS(O)_a wherein a is 0, C₁₋₄alkoxy, amino, carbocyclyl, heterocyclyl or mercapto; wherein R²⁰ may be independently optionally substituted on carbon
20 by one or more R²²;
R²² is selected from hydroxy; and
R²³ is carboxy.

In another further aspect of the invention, more preferably R¹³ is hydrogen, methyl, ethyl, butyl or phenyl or R²³; wherein R¹³ is optionally substituted by one or more substituents
25 selected from R²⁰; wherein

R²⁰ is hydroxy, methylthio, methoxy, amino, imidazolyl or mercapto; wherein R²⁰ may be independently optionally substituted on carbon by one or more R²²;

R²² is selected from hydroxy; and
R²³ is carboxy.
30 In another further aspect of the invention, particularly R¹³ is hydrogen, carboxy, hydroxymethyl, mercaptomethyl, methoxymethyl, methylthiomethyl, 2-methylthioethyl, 4-aminobutyl, 4-hydroxybenzyl, imidazol-5-ylmethyl or phenyl.

In an other aspect more particularly R¹³ is methylthiomethyl, methylsulphinylmethyl or methylsulphonylmethyl.

Preferably R¹⁴ is hydrogen.

In another aspect of the invention, preferably R¹⁴ is selected from hydrogen, C₁₋₄alkyl or carbocyclyl; wherein said C₁₋₄alkyl or carbocyclyl may be optionally substituted by one or more substituents selected from R²⁰; and

R²⁰ is hydroxy.

In another aspect of the invention, more preferably R¹⁴ is selected from hydrogen, methyl or phenyl; wherein said methyl or phenyl may be optionally substituted by one or more substituents selected from R²⁰; and

R²⁰ is hydroxy.

In another aspect of the invention, particularly R¹⁴ is hydrogen, phenyl or hydroxymethyl.

Particularly R¹⁵ is carboxy or sulpho.

15 In one aspect of the invention, more particularly R¹⁵ is carboxy.

In another aspect of the invention, more particularly R¹⁵ is sulpho.

Preferably R¹⁵ is carboxy, sulpho, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₄alkyl.

More preferably R¹⁵ is carboxy, sulpho, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), 20 -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from methyl or ethyl.

Preferably R¹⁵ is carboxy, sulpho, -P(O)(OEt)(OEt), -P(O)(OH)(OEt), -P(O)(OH)(Me) or -P(O)(OEt)(Me).

Preferably R¹⁵ is carboxy, sulpho, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), 25 -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₄alkyl or R¹⁵ is a group of formula (IC) (as depicted above).

More preferably R¹⁵ is carboxy, sulpho, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from methyl or ethyl or R¹⁵ is a group of formula (IC) (as depicted above).

30 Preferably R¹⁵ is carboxy, sulpho, phosphono, -P(O)(OEt)(OEt), -P(O)(O*t*-Bu)(O*t*-Bu), -P(O)(OH)(OEt), -P(O)(OH)(Me) or -P(O)(OEt)(Me) or R¹⁵ is a group of formula (IC) (as depicted above).

In one aspect of the invention, preferably R¹⁵ is a group of formula (IC) (as depicted above).

In another aspect of the invention, preferably R¹⁵ is not a group of formula (IC) (as depicted above).

5 In one aspect of the invention, preferably R¹⁵ is carboxy.

In another aspect of the invention, preferably R¹⁵ is sulpho.

In another aspect of the invention, preferably R¹⁵ is -P(O)(OH)(OEt).

In another aspect of the invention, preferably R¹⁵ is -P(O)(OH)(Me).

In another aspect of the invention, preferably R¹⁵ is -P(O)(OEt)(Me).

10 In one aspect of the invention, preferably R²⁴ is hydrogen.

In another aspect of the invention, preferably R²⁴ is C₁₋₄alkyl.

Preferably R²⁵ is hydrogen.

Preferably R²⁶ is carboxy.

Preferably p is 1 or 2; wherein the values of R¹³ may be the same or different.

15 In one aspect of the invention, more preferably p is 1.

In another aspect of the invention, more preferably p is 2; wherein the values of R¹³ may be the same or different.

In a further aspect of the invention, more preferably p is 3; wherein the values of R¹³ may be the same or different.

20 In one aspect of the invention, preferably q is 0.

In a further aspect of the invention, preferably q is 1.

In one aspect of the invention, preferably r is 0.

In one aspect of the invention, more preferably r is 1.

25 In another aspect of the invention, more preferably r is 2; wherein the values of R¹⁴ may be the same or different.

In a further aspect of the invention, more preferably r is 3; wherein the values of R¹⁴ may be the same or different.

Preferably m is 0.

In another aspect of the invention, preferably m is 0 or 1.

30 Preferably n is 1.

In another aspect of the invention, preferably n is 1 or 2.

Preferably z is 1.

The group of formula (IA') wherein R⁷ is hydrogen, methyl or phenyl, n is 1, Ring A is phenyl, thienyl or indolyl; wherein Ring A is optionally substituted by one or more substituents selected from halo, hydroxy or trifluoromethyl, m is 0 and R⁹ is carboxy, -P(O)(OH)(OR^c) or a group of formula (IB).

5 The group of formula (IA) wherein:

D is -O- or -S-;

Ring A is phenyl, thienyl or indolyl; wherein Ring A is optionally substituted by one or more substituents selected from halo, hydroxy, methoxy or trifluoromethyl;

R⁷ is hydrogen, methyl or phenyl;

10 R⁸ is hydrogen or methyl;

R⁹ is hydrogen or methyl;

R¹⁰ is hydrogen;

m is 0-2 wherein the values of R¹⁰ may be the same or different; and

R¹¹ is carboxy, -P(O)(OH)(OEt) or a group of formula (IB) (as depicted in claim 1);

15 The group of formula (IB') wherein R¹⁰ is hydrogen, hydroxymethyl or phenyl, p is 1 or 2; wherein the values of R¹⁰ may be the same or different and R¹¹ is carboxy or sulpho.

The group of formula (IB) wherein:

R¹² is hydrogen or methyl;

R¹³ is hydrogen, methyl, ethyl, butyl or phenyl or R²³; wherein R¹³ is optionally

20 substituted by one or more substituents selected from R²⁰; R²⁰ is hydroxy, methylthio, methoxy, amino, imidazolyl or mercapto; wherein R²⁰ may be independently optionally substituted on carbon by one or more hydroxy; R²³ is carboxy;

X is -NH- or -NHC(O)-;

25 R¹⁴ is selected from hydrogen, methyl or phenyl; wherein said methyl or phenyl may be optionally substituted by one or more substituents selected from hydroxy;

R¹⁵ is carboxy, sulpho, phosphono, -P(O)(ORE)(OR^f), -P(O)(OH)(ORE), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from methyl or ethyl or R¹⁵ is a group of formula (IC) (as depicted in claim 1);

p is 1-3 wherein the values of R¹³ may be the same or different;

30 q is 0-1; and

r is 0-3 wherein the values of R¹⁴ may be the same or different;

The group of formula (IC) wherein

R²⁴ is hydrogen;

R^{25} is hydrogen;

R^{26} is carboxy; and

z is 1;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug

5 thereof.

Therefore in a further aspect of the invention, there is provided a compound of formula (I') as depicted above wherein:

R^1 and R^2 are independently selected from ethyl or butyl;

R^3 and R^6 are hydrogen;

10 R^4 is selected from halo, C_{1-4} alkoxy or C_{1-4} alkylS(O)_a wherein a is 0 to 2; wherein that R^4 may be optionally substituted on carbon by one or more R^{16} ; wherein R^{16} is independently selected from hydroxy and N,N -(C_{1-4} alkyl)₂amino;

R^5 is a group of formula (IA');

15 Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R^{17} ; wherein

R^{17} is selected from halo, hydroxy or C_{1-4} alkyl; wherein R^{17} may be optionally substituted on carbon by one or more R^{21} ; wherein

R^{21} is selected from halo;

R^7 is hydrogen, C_{1-4} alkyl or carbocyclyl;

20 R^{11} is carboxy, $-P(O)(OH)(OR^C)$ or a group of formula (IB') (as depicted above);

R^{13} is hydrogen, C_{1-4} alkyl or carbocyclyl; wherein R^{13} is optionally substituted by one or more substituents selected from R^{20} ; wherein

R^{20} is hydroxy;

R^{15} is carboxy or sulpho;

25 p is 1 or 2; wherein the values of R^{13} may be the same or different;

m is 0; and

n is 1;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore in an additional aspect of the invention, there is provided a compound of formula (I') as depicted above wherein:

R^1 and R^2 are both butyl or one of R^1 and R^2 is ethyl and the other is butyl;

R^4 is methylthio;

R^5 is a group of formula (IA') (as depicted above);

R³ and **R⁶** are hydrogen;

Ring A is phenyl;

R⁷ is hydrogen;

R¹¹ is a group of formula (IB') (as depicted above);

5 **R¹³** is hydrogen or hydroxymethyl;

R¹⁵ is carboxy or sulpho;

p is 1 or 2; wherein the values of **R¹³** may be the same or different;

m is 0;

n is 1;

10. or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore in an additional further aspect of the invention, there is provided a compound of formula (I'') as depicted above wherein:

R¹ and **R²** are independently selected from ethyl or butyl;

R³ and **R⁶** are hydrogen;

15. **R⁴** is selected from halo, C₁₋₄alkoxy or C₁₋₄alkylS(O)_a wherein a is 0 to 2; wherein that **R⁴** may be optionally substituted on carbon by one or more **R¹⁶**; wherein **R¹⁶** is independently selected from hydroxy and *N,N*-(C₁₋₄alkyl)₂amino;

R⁵ is a group of formula (IA'');

20 **Ring A** is aryl or heteroaryl; wherein **Ring A** is optionally substituted by one or more substituents selected from **R¹⁷**;

R⁷ is hydrogen, C₁₋₄alkyl or carbocyclyl;

R⁸ is hydrogen or methyl;

R⁹ is hydrogen or methyl;

R¹¹ is carboxy, -P(O)(OH)(OR^c) or a group of formula (IB'') (as depicted above);

25 **X** is -NH- or -NHC(O)-;

R¹² is hydrogen or methyl;

R¹³ is hydrogen, C₁₋₄alkyl or carbocyclyl; wherein **R¹³** is optionally substituted by one or more substituents selected from **R²⁰**;

R¹⁴ is hydrogen;

30 **R¹⁵** is carboxy or sulpho;

R¹⁷ is selected from halo, hydroxy, C₁₋₄alkyl or C₁₋₄alkoxy; wherein **R¹⁷** may be optionally substituted on carbon by one or more **R²¹**;

R^{20} is hydroxy, carboxy, carbocyclyl or amino; wherein R^{20} may be optionally substituted on carbon by one or more R^{22} ;

R^{21} is selected from halo;

R^{22} is hydroxy;

5 p is 1 - 3; wherein the values of R^{13} may be the same or different.

q is 0 - 1;

r is 0 - 3; wherein the values of R^{14} may be the same or different; and wherein if q is 1, r is not 0;

m is 0 - 2; and

10 n is 1 - 3;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore in another additional further aspect of the invention, there is provided a compound of formula (I) as depicted above wherein:

R^v and R^w are both hydrogen;

15 R^1 and R^2 are independently selected from C_{1-4} alkyl;

R^x and R^y are both hydrogen;

R^z is selected from halo, amino, C_{1-6} alkyl, C_{1-6} alkoxycarbonylamino or

$N'-(C_{1-6}$ alkyl)ureido;

v is 0 or 1;

20 R^3 and R^6 are hydrogen;

 one of R^4 and R^5 is a group of formula (IA) (as depicted above) and the other is selected from hydrogen, halo, C_{1-4} alkoxy or C_{1-4} alkylS(O)_a wherein a is 0 to 2; wherein that R^4 or R^5 may be optionally substituted on carbon by one or more R^{16} ; wherein R^{16} is independently selected from hydroxy, carboxy and $N,N-(C_{1-4}$ alkyl)₂amino;

25 D is -O- or -S-;

R^7 is hydrogen, methyl or phenyl;

R^8 is hydrogen or methyl;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R^{17} ; wherein R^{17} is selected from halo, hydroxy, C_{1-4} alkyl or

30 C_{1-4} alkoxy; wherein R^{17} may be optionally substituted on carbon by one or more R^{21} ; wherein R^{21} is selected from halo;

R^9 is hydrogen or methyl;

R^{10} is hydrogen;

R¹¹ is carboxy, -P(O)(OH)(OR^c) wherein R^c is selected from C₁₋₄alkyl or a group of formula (IB) (as depicted above);

R¹² is hydrogen or methyl;

X is -NH- or -NHC(O)-;

5 **R¹³** is hydrogen, C₁₋₄alkyl, carbocyclyl or R²³; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein R²⁰ is hydroxy, C₁₋₄alkylS(O)_a wherein a is 0, C₁₋₄alkoxy, amino, carbocyclyl, heterocyclyl or mercapto; wherein R²⁰ may be independently optionally substituted on carbon by one or more R²²; R²² is selected from hydroxy; and R²³ is carboxy;

10 **R¹⁴** is selected from hydrogen, C₁₋₄alkyl or carbocyclyl; wherein said C₁₋₄alkyl or carbocyclyl may be optionally substituted by one or more substituents selected from R²⁰; and R²⁰ is hydroxy;

R¹⁵ is carboxy, sulpho, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from

15 C₁₋₄alkyl or R¹⁵ is a group of formula (IC) (as depicted above);

R²⁴ is hydrogen;

R²⁵ is hydrogen;

R²⁶ is carboxy;

p is 1-3; wherein the values of R¹³ may be the same or different;

20 **q** is 0-1;

r is 0-3; wherein the values of R¹⁴ may be the same or different;

m is 0-2; wherein the values of R¹⁰ may be the same or different;

n is 1-2; wherein the values of R⁷ may be the same or different;

z is 0-1; wherein the values of R²⁵ may be the same or different;

25 or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

In another aspect of the invention, preferred compounds of the invention are any one of the Examples or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

30 In one aspect of the invention, there is provided a compound of formula (I) selected from Examples 8, 9, 46, 56, 59, 60, 61, 62, 66 and 69 or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

In another aspect of the invention, there is provided a compound of formula (I) which is Example 73, 74, 95, 96, 97, 98, 99 and 100 or a pharmaceutically acceptable salt, solvate,

solvate of such a salt or a prodrug thereof.

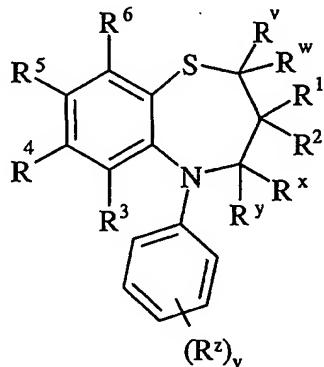
In another aspect of the invention, preferred compounds of the invention are any one of Examples 43, 50, 51 and 52 or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

5 In another additional aspect of the invention, preferred compounds of the invention are any one of Examples 43, 46, 50, 51, 56, 58, 59, 61, 62, 63, 69, 81, 83, 85, 94, 97, 98, 108, 109, 110, 111 or 117.

Preferred aspects of the invention are those which relate to the compound of formula (I) or a pharmaceutically acceptable salt thereof.

10 Another aspect of the present invention provides a process for preparing a compound of formula (I) or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof which process (wherein variable groups are, unless otherwise specified, as defined in formula (I)) comprises of:

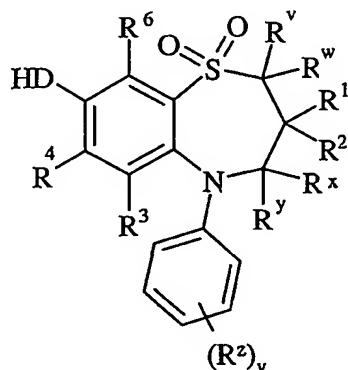
Process 1): oxidising a benzothiazepine of formula (II):



15

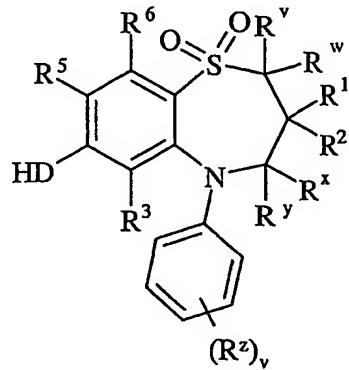
(II);

Process 2): for compounds of formula (I) wherein D is -O-, -NR^a or -S-; reacting a compound of formula (IIIa) or (IIIb):



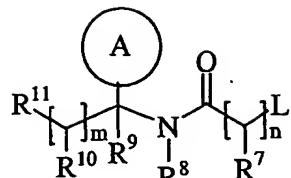
20

(IIIa)



(IIIb)

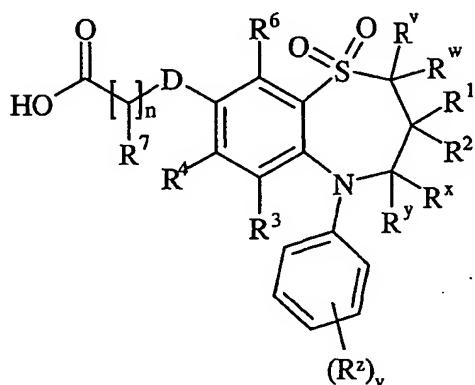
with a compound of formula (IV):



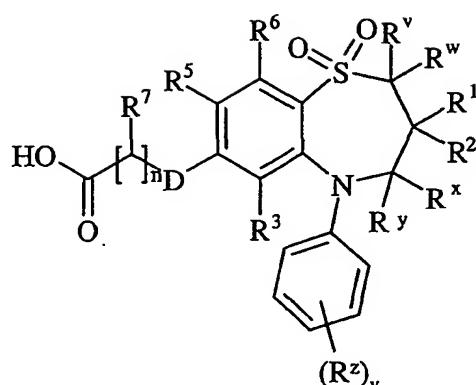
(IV)

wherein L is a displaceable group;

5 Process 3): reacting an acid of formula (Va) or (Vb):

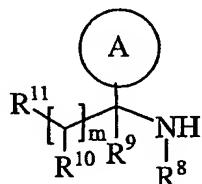


(Va)



(Vb)

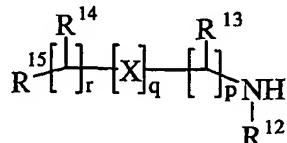
or an activated derivative thereof; with an amine of formula (VI):



10

(VI);

Process 4): for compounds of formula (I) wherein R¹¹ is a group of formula (IB); reacting a compound of formula (I) wherein R¹¹ is carboxy with an amine of formula (VII):

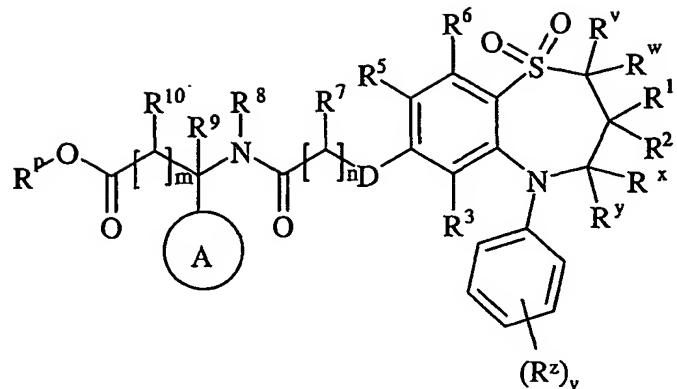


15

(VII)

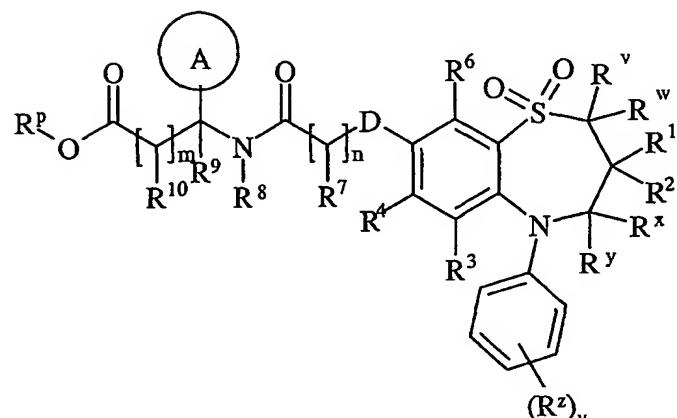
Process 5): for compounds of formula (I) wherein R¹¹ is carboxy; deprotecting a compound of formula (VIIIa):

- 29 -



(VIIIa)

or (VIIIb):



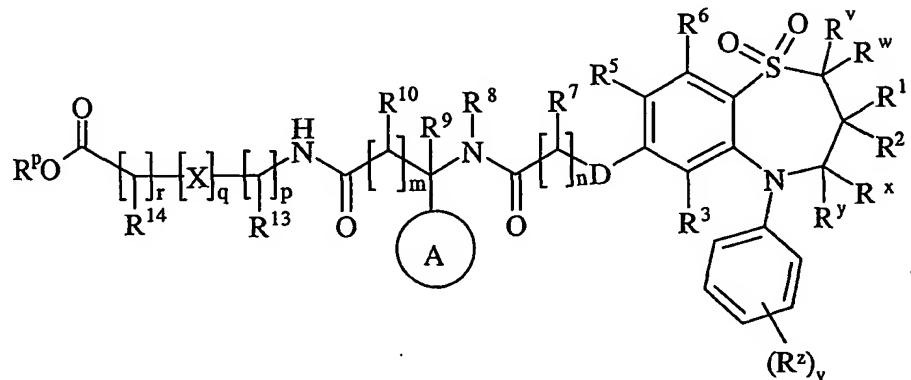
5

(VIIIb)

wherein R^p is C₁₋₄alkyl;

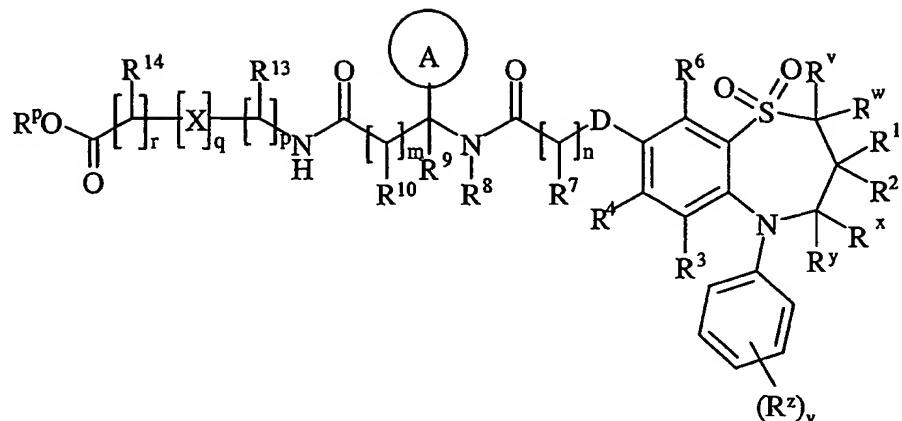
Process 6): for compounds of formula (I) wherein R¹¹ is a group of formula (IB) and R¹⁵ is carboxy; deprotecting a compound of formula (IXa):

10



(IXa)

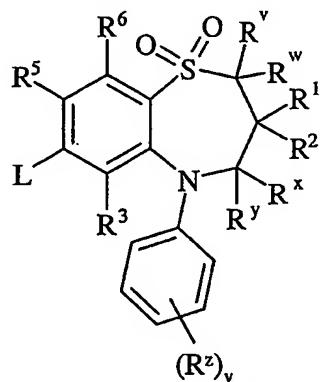
or (IXb):



(IXb)

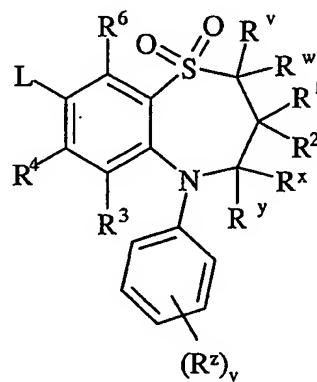
5 wherein R^p is C₁₋₄alkyl;

Process 7) for compounds of formula (I) wherein one of R^4 and R^5 are independently selected from C₁₋₄alkylthio optionally substituted on carbon by one or more R^{16} ; reacting a compound of formula (Xa) or (Xb):



10

(Xa)



(Xb)

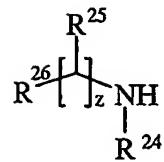
wherein L is a displaceable group; with a thiol of formula (XI):



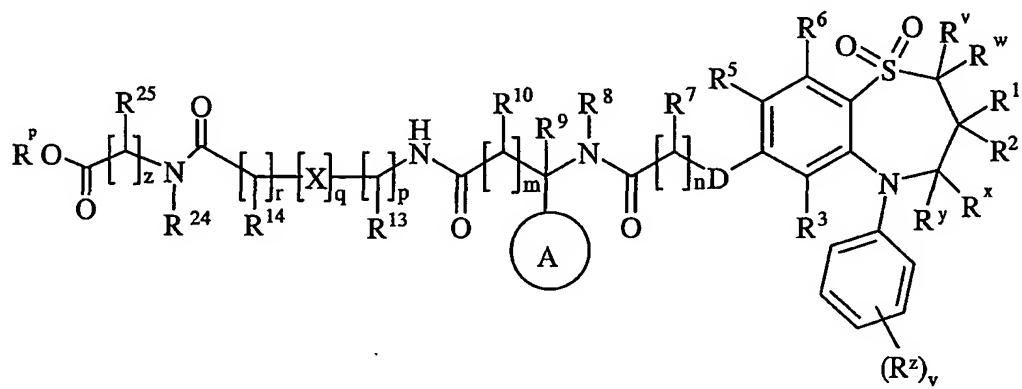
(XI)

15 wherein R^y is C₁₋₄alkylthio optionally substituted on carbon by one or more R^{16} ;

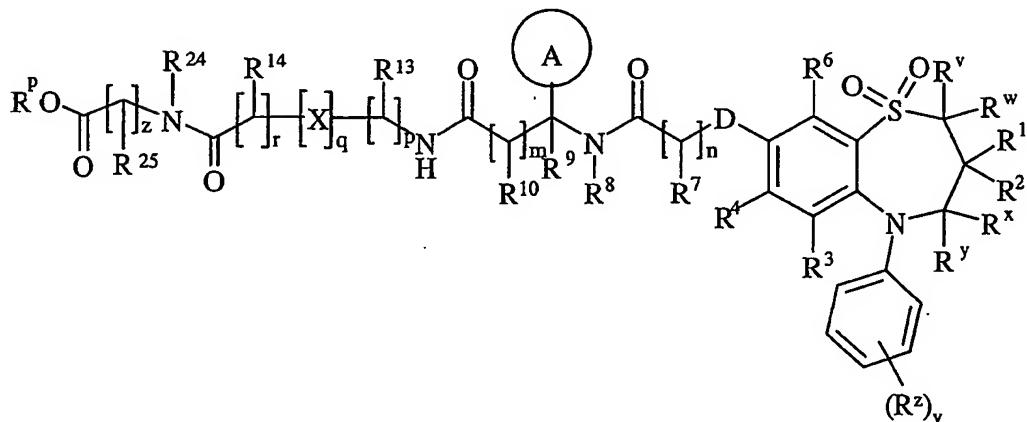
Process 8) for compounds of formula (I) wherein R^{15} is a group of formula (IC) reacting a compound of formula (IXa) or (IXb) wherein R^p is hydrogen with a compound of formula (XII):



Process 9): for compounds of formula (I) wherein R¹¹ is a group of formula (IB) and R¹⁵ is a group of formula (IC) and R²⁶ is carboxy; deprotecting a compound of formula (XIIIa):

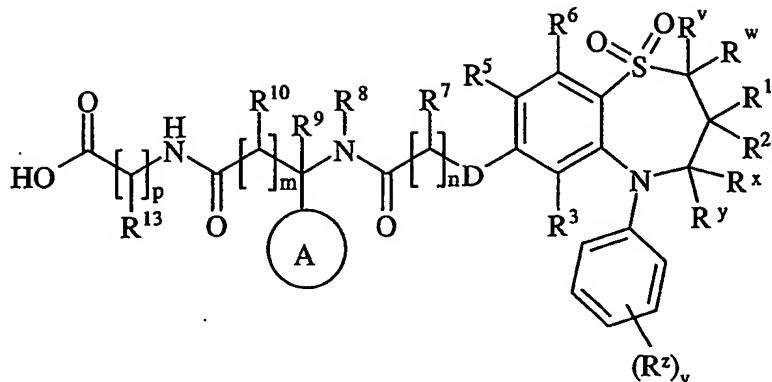


or (XIIIb):



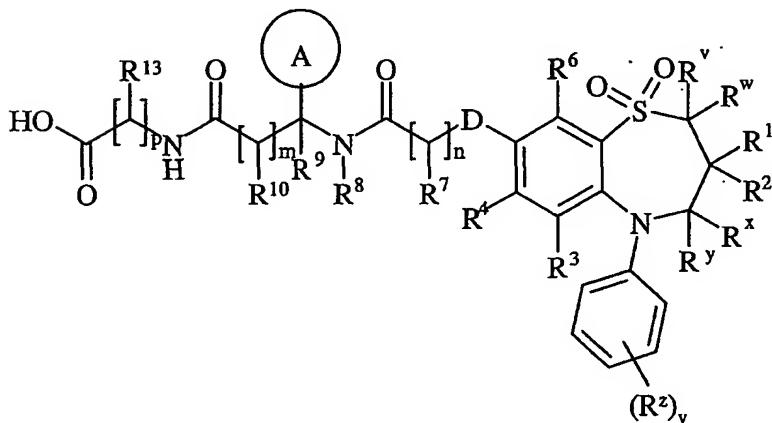
10 and R^p is C₁₋₄alkyl;

Process 10): for compounds of formula (I) wherein X is -N(R^q)C(O)-; reacting a compound of formula (XIVa):



(XIVa)

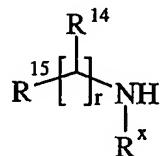
or (XIVb):



5

(XIVb)

with a compound of formula (XV):



(XV)

and thereafter if necessary or desirable:

10 i) converting a compound of the formula (I) into another compound of the formula (I);
ii) removing any protecting groups;
iii) forming a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug.

The skilled person will also appreciate that similar processes corresponding to the above processes can also be used to prepare compounds of formula (I') and compounds of
15 formula (I'') wherein the definitions of the variable groups may differ.

L is a displaceable group, suitable values for L are for Example, a halogeno or

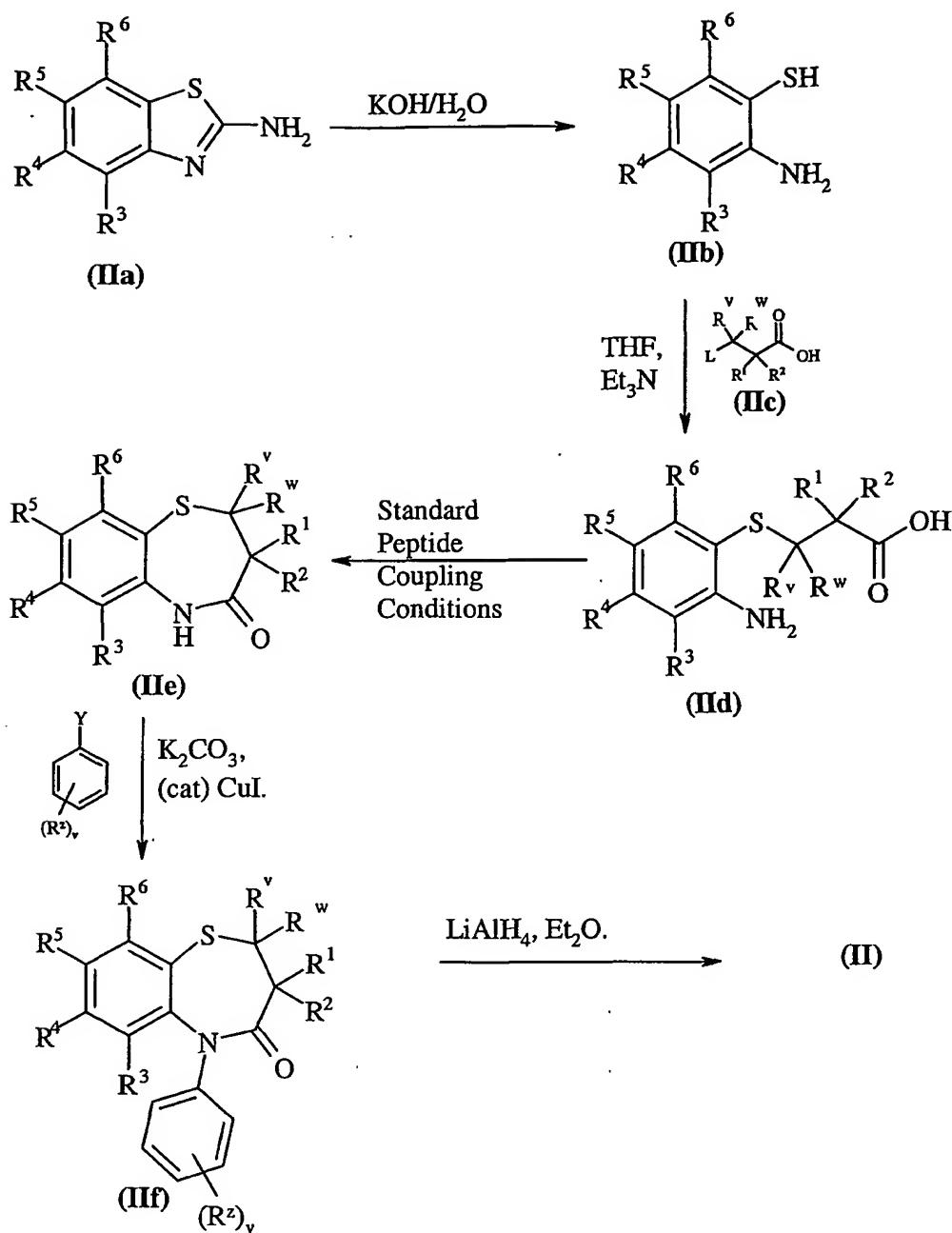
sulphonyloxy group, for Example a chloro, bromo, methanesulphonyloxy or toluene-4-sulphonyloxy group.

R^P is C₁₋₄alkyl. Preferably R^P is methyl or ethyl. More preferably R^P is methyl.

Specific reaction conditions for the above reactions are as follows.

5 *Process 1*): Benzothiazepines of formula (II) may be oxidised under standard sulphur oxidation conditions; for Example using hydrogen peroxide and trifluoroacetic acid at a temperature in the range of 0°C to reflux, preferably at or near room temperature.

Compounds of formula (II) may be prepared according to Scheme I for compounds of formula (I) wherein R^x and R^y are hydrogen. The skilled man will appreciate that where R^x and R^y are not both hydrogen the following synthetic route needs to be manipulated using procedures known to the skilled person.

*Scheme 1*

wherein L is a displaceable group as defined above, and Y is a displaceable group, for
 5 Example halo.

Compounds of formula (IIa) and (IIc) are commercially available compounds, or they are known in the literature, or they are prepared by standard processes known in the art.

Process 2): Alcohols of formula (IIIa) or (IIIb) may be reacted with compounds of formula (IV) in the presence of a base for Example an inorganic base such as sodium carbonate, or an

organic base such as Hunigs base, in the presence of a suitable solvent such as acetonitrile, dichloromethane or tetrahydrofuran at a temperature in the range of 0°C to reflux, preferably at or near reflux.

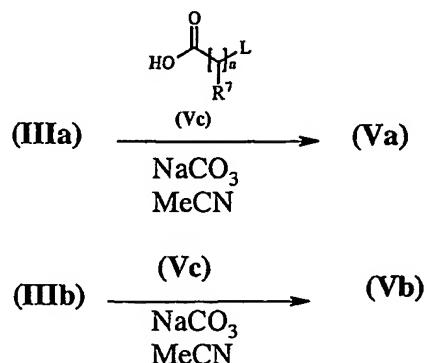
Compounds of formula (IIIa) or (IIIb) may be prepared in a similar manner to 5 compounds of formula (II) (but wherein R⁴ or R⁵ is hydroxy) followed by the oxidation step of Process 1).

Compounds of formula (IV) are commercially available compounds, or they are known in the literature, or they are prepared by standard processes known in the art.

Process 3), Process 4, Process 8) and Process 10): Acids and amines may be coupled 10 together in the presence of a suitable coupling reagent. Standard peptide coupling reagents known in the art can be employed as suitable coupling reagents, or for Example carbonyldiimidazole and dicyclohexyl-carbodiimide, optionally in the presence of a catalyst such as dimethylaminopyridine or 4-pyrrolidinopyridine, optionally in the presence of a base for Example triethylamine, pyridine, or 2,6-di-*alkyl*-pyridines such as 2,6-lutidine or 15 2,6-di-*tert*-butylpyridine. Suitable solvents include dimethylacetamide, dichloromethane, benzene, tetrahydrofuran and dimethylformamide. The coupling reaction may conveniently be performed at a temperature in the range of -40 to 40°C.

Suitable activated acid derivatives include acid chlorides, for Example acid chlorides, and active esters, for Example pentafluorophenyl esters. The reaction of these types of 20 compounds with amines is well known in the art, for Example they may be reacted in the presence of a base, such as those described above, and in a suitable solvent, such as those described above. The reaction may conveniently be performed at a temperature in the range of -40 to 40°C.

Compounds of formula (Va) or (Vb) wherein D is -O-, -NR^a- or -S- may be prepared 25 according to Scheme 2:

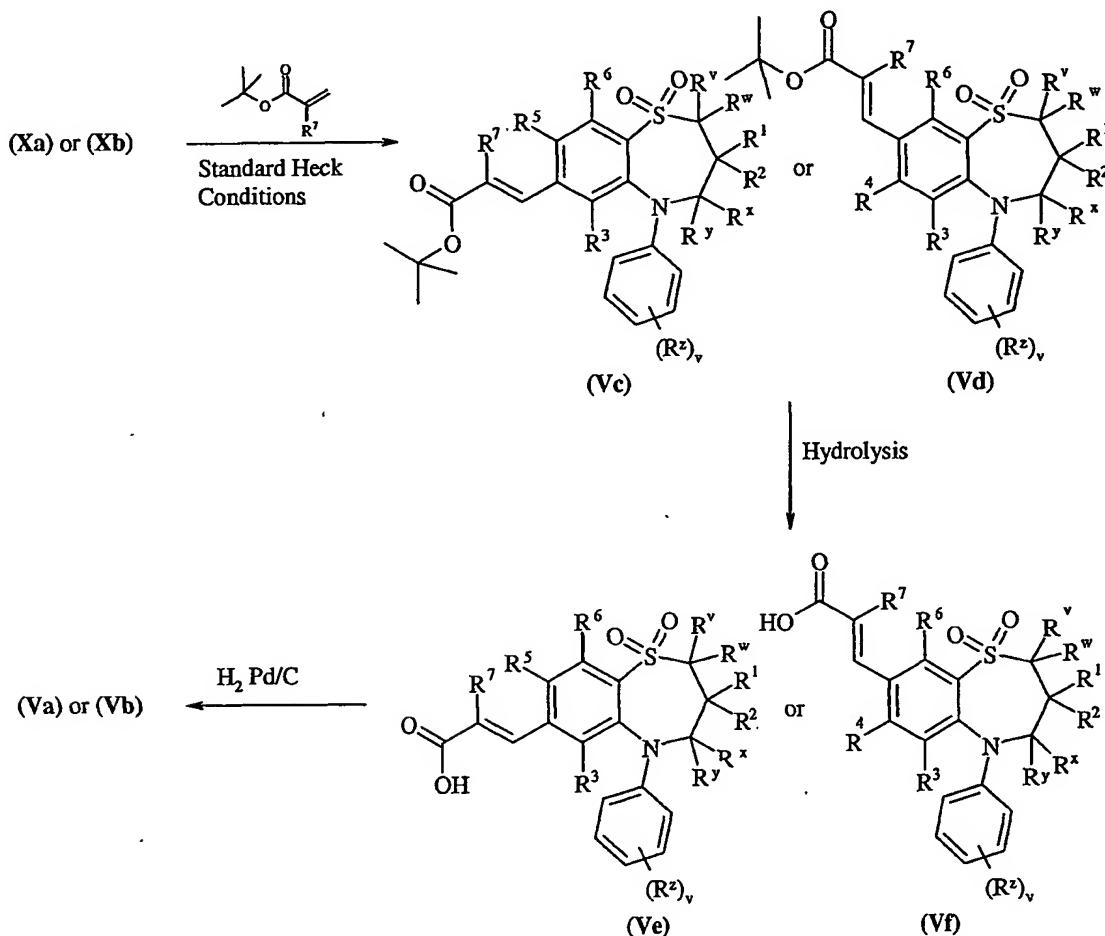


Scheme 2

wherein L is a displaceable group as defined above.

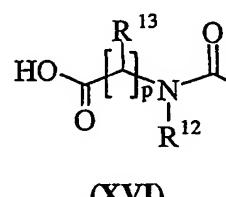
Compounds of formula (Va) and (Vb) where D is -SO- or -SO₂- may be prepared by oxidising the resulting compounds of formula (Va) and (Vb) from *Scheme 2* where D is -S-.

Compounds of formula (Va) or (Vb) wherein D is -CH₂- may be prepared according to *Scheme 3*.



Scheme 3

Compounds of formula (XIVa) or (XIVa) may be prepared by any of the processes described herein where R¹¹ is a group of formula (IB) but wherein (IB) is a group of formula 10 (XVI):



Compounds of formula (Vc), (VI), (VII), (XII), (XV) and (XVI) are commercially available compounds, or they are known in the literature, or they are prepared by standard processes known in the art.

Process 5), Process 6) and Process 9): Esters of formula (VIIIa), (VIIIb), (IXa), (IXb)

5 (XIIIa) and (XIIIb) may be deprotected under standard conditions such as those described below, for Example they may be deprotected with sodium hydroxide in methanol at room temperature.

Esters of formula (VIIIa), (VIIIb), (IXa), (IXb) (XIIIa) and (XIIIb) may be prepared by any of the procedures above for the preparation of compounds of formula (I), but wherein
10 R¹¹, R¹⁵ or R²⁶ is C₁₋₄alkoxycarbonyl.

Process 7): Compounds of formula (Xa) and (Xb) may be reacted with thiols of formula (XI) in the presence of base, for Example an inorganic base such as sodium carbonate or an organic base such as Hunigs base, in the presence of a suitable solvent such as DMF or THF at a temperature in the range of 0°C to reflux.

15 Compounds of formula (Xa) and (Xb) may be prepared by any of the procedures above for the preparation of compounds of formula (I), but wherein one of R⁴ and R⁵ is L.

Compounds of formula (XI) are commercially available compounds, or they are known in the literature, or they are prepared by standard processes known in the art.

It will be appreciated that certain of the various ring substituents in the compounds of
20 the present invention may be introduced by standard aromatic substitution reactions or generated by conventional functional group modifications either prior to or immediately following the processes mentioned above, and as such are included in the process aspect of the invention. Such reactions and modifications include, for example, introduction of a substituent by means of an aromatic substitution reaction, reduction of substituents, alkylation
25 of substituents and oxidation of substituents. The reagents and reaction conditions for such procedures are well known in the chemical art. Particular examples of aromatic substitution reactions include the introduction of a nitro group using concentrated nitric acid, the introduction of an acyl group using, for example, an acyl halide and Lewis acid (such as aluminium trichloride) under Friedel Crafts conditions; the introduction of an alkyl group
30 using an alkyl halide and Lewis acid (such as aluminium trichloride) under Friedel Crafts conditions; and the introduction of a halogeno group. Particular examples of modifications include the reduction of a nitro group to an amino group by for example, catalytic

hydrogenation with a nickel catalyst or treatment with iron in the presence of hydrochloric acid with heating; oxidation of alkylthio to alkylsulphinyloxy or alkylsulphonyloxy.

It will also be appreciated that in some of the reactions mentioned herein it may be necessary/desirable to protect any sensitive groups in the compounds. The instances where 5 protection is necessary or desirable and suitable methods for protection are known to those skilled in the art. Conventional protecting groups may be used in accordance with standard practice (for illustration see T.W. Green, Protective Groups in Organic Synthesis, John Wiley and Sons, 1991). Thus, if reactants include groups such as amino, carboxy or hydroxy it may be desirable to protect the group in some of the reactions mentioned herein.

10 A suitable protecting group for an amino or alkylamino group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an alkoxycarbonyl group, for example a methoxycarbonyl, ethoxycarbonyl or *t*-butoxycarbonyl group, an arylmethoxycarbonyl group, for example benzyloxycarbonyl, or an aroyl group, for example benzoyl. The deprotection conditions for the above protecting groups necessarily vary with the choice of protecting

15 group. Thus, for example, an acyl group such as an alkanoyl or alkoxycarbonyl group or an aroyl group may be removed for example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an acyl group such as a *t*-butoxycarbonyl group may be removed, for example, by treatment with a suitable acid as hydrochloric, sulphuric or phosphoric acid or trifluoroacetic acid and an

20 arylmethoxycarbonyl group such as a benzyloxycarbonyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon, or by treatment with a Lewis acid for example boron tris(trifluoroacetate). A suitable alternative protecting group for a primary amino group is, for example, a phthaloyl group which may be removed by treatment with an alkylamine, for example dimethylaminopropylamine, or with hydrazine.

25 A suitable protecting group for a hydroxy group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an aroyl group, for example benzoyl, or an arylmethyl group, for example benzyl. The deprotection conditions for the above protecting groups will necessarily vary with the choice of protecting group. Thus, for example, an acyl group such as an alkanoyl or an aroyl group may be removed, for example, by hydrolysis with 30 a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an arylmethyl group such as a benzyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

A suitable protecting group for a carboxy group is, for example, an esterifying group, for example a methyl or an ethyl group which may be removed, for example, by hydrolysis with a base such as sodium hydroxide, or for example a *t*-butyl group which may be removed, for example, by treatment with an acid, for example an organic acid such as trifluoroacetic acid, or for example a benzyl group which may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

The protecting groups may be removed at any convenient stage in the synthesis using conventional techniques well known in the chemical art.

As stated hereinbefore the compounds defined in the present invention possess IBAT inhibitory activity. These properties may be assessed, for example, using an *in vitro* test assay for studying the effect on bile acid uptake in IBAT-transfected cells (Smith L., Price-Jones M. J., Hugnes K. T. and Jones N. R. A.; J Biomolecular Screening, 3, 227-230) or *in vivo* by studying the effect on radiolabelled bile acid absorption in mice/rats (Lewis M. C., Brieaddy L. E. and Root C., J., J Lip Res 1995, 36, 1098-1105).

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in association with a pharmaceutically-acceptable diluent or carrier.

The composition may be in a form suitable for oral administration, for example as a tablet or capsule, for parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion) as a sterile solution, suspension or emulsion, for topical administration as an ointment or cream or for rectal administration as a suppository.

In general the above compositions may be prepared in a conventional manner using conventional excipients.

The compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, will normally be administered to a warm-blooded animal at a unit dose within the range 5-5000 mg per square meter body area of the animal, i.e. approximately 0.1-100 mg/kg or 0.01-50 mg/kg, and this normally provides a therapeutically-effective dose. A unit dose form such as a tablet or capsule will usually contain, for example 1-250 mg of active ingredient. Preferably a daily dose in the range of 1-50 mg/kg is employed. In another aspect a daily dose in the range of 0.02-20 mg/kg is employed. However the daily dose will necessarily be varied depending upon the host treated, the particular route of administration, and the severity of the illness being treated.

Accordingly the optimum dosage may be determined by the practitioner who is treating any particular patient.

According to a further aspect of the present invention there is provided a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore for use in a method of prophylactic or therapeutic treatment of a warm-blooded animal, such as man.

We have found that the compounds defined in the present invention, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, are effective IBAT inhibitors, and accordingly have value in the treatment of disease states associated with hyperlipidaemic conditions.

Thus according to this aspect of the invention there is provided a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore for use as a medicament.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in the manufacture of a medicament for use in the treatment of hyperlipidaemic conditions in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in the manufacture of a medicament for use in the treatment of dyslipidemic conditions and disorders such as hyperlipidaemia, hypertriglyceridemia, hyperbetalipoproteinemia (high LDL), hyperprebetalipoproteinemia (high VLDL), hyperchylomicronemia, hypolipoproteinemia, hypercholesterolemia, hyperlipoproteinemia and hypoalphalipoproteinemia (low HDL) in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in the manufacture of a medicament for use in the treatment of different clinical conditions such as atherosclerosis, arteriosclerosis, arrhythmia,

hyper-thrombotic conditions, vascular dysfunction, endothelial dysfunction, heart failure, coronary heart diseases, cardiovascular diseases, myocardial infarction, angina pectoris, peripheral vascular diseases, inflammation of cardiovascular tissues such as heart, valves, vasculature, arteries and veins, aneurisms, stenosis, restenosis, vascular plaques, vascular fatty streaks, leukocyte, monocytes and/or macrophage infiltrate, intimal thickening, medial thinning, infectious and surgical trauma and vascular thrombosis, stroke and transient ischaemic attacks in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in the manufacture of a medicament for use in the treatment of atherosclerosis, coronary heart diseases, myocardial infarction, angina pectoris, peripheral vascular diseases, stroke and transient ischaemic attacks in a warm-blooded animal, such as man.

According to a further feature of this aspect of the invention there is provided a method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

According to a further feature of this aspect of the invention there is provided a method of treating hyperlipidemic conditions in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

According to a further feature of this aspect of the invention there is provided a method of treating dyslipidemic conditions and disorders such as hyperlipidaemia, hypertriglyceridemia, hyperbetalipoproteinemia (high LDL), hyperprebetalipoproteinemia (high VLDL), hyperchylomicronemia, hypolipoproteinemia, hypercholesterolemia, hyperlipoproteinemia and hypoalphalipoproteinemia (low HDL) in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

According to a further feature of this aspect of the invention there is provided a method of treating different clinical conditions such as atherosclerosis, arteriosclerosis,

arrhythmia, hyper-thrombotic conditions, vascular dysfunction, endothelial dysfunction, heart failure, coronary heart diseases, cardiovascular diseases, myocardial infarction, angina pectoris, peripheral vascular diseases, inflammation of cardiovascular tissues such as heart, valves, vasculature, arteries and veins, aneurisms, stenosis, restenosis, vascular plaques,

5 vascular fatty streaks, leukocyte, monocytes and/or macrophage infiltrate, intimal thickening, medial thinning, infectious and surgical trauma and vascular thrombosis, stroke and transient ischaemic attacks in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

10 According to a further feature of this aspect of the invention there is provided a method of treating atherosclerosis, coronary heart diseases, myocardial infarction, angina pectoris, peripheral vascular diseases, stroke and transient ischaemic attacks in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

15 There is evidence that an IBAT inhibitor might potentially be useful in the treatment and/or prevention of gallstones. According to a further feature of this aspect of the invention there is provided a method of treating and / or preventing gallstones in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal

20 an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

The size of the dose required for the therapeutic or prophylactic treatment will necessarily be varied depending on the host treated, the route of administration and the severity of the illness being treated. A unit dose in the range, for example, 1-100 mg/kg,

25 preferably 1-50 mg/kg is envisaged.

The IBAT inhibitory activity defined hereinbefore may be applied as a sole therapy or may involve, in addition to a compound of the invention, one or more other substances and/or treatments. Such conjoint treatment may be achieved by way of the simultaneous, sequential or separate administration of the individual components of the treatment. According to this

30 aspect of the invention there is provided a pharmaceutical product comprising a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore and an additional IBAT inhibitory substance as

defined hereinbefore and an additional hypolipidaemic agent for the conjoint treatment of hyperlipidaemia.

In another aspect of the invention, the compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, may be administered in association with an HMG Co-A reductase inhibitor, or pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof. Suitable HMG Co-A reductase inhibitors, pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof are statins well known in the art. Particular statins are fluvastatin, lovastatin, pravastatin, simvastatin, atorvastatin, cerivastatin, bervastatin, dalvastatin, mevastatin and (E)-7-[4-(4-fluorophenyl)-6-isopropyl-2-[methyl(methylsulphonyl)amino]pyrimidin-5-yl](3R,5S)-3,5-dihydroxyhept-6-enoic acid (rosuvastatin), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof. A particular statin is atorvastatin, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof. A more particular statin is atorvastatin calcium salt. A further particular statin is (E)-7-[4-(4-fluorophenyl)-6-isopropyl-2-[methyl(methylsulphonyl)amino]pyrimidin-5-yl](3R,5S)-3,5-dihydroxyhept-6-enoic acid (rosuvastatin), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof. A preferable particular statin is rosuvastatin calcium salt.

In an additional aspect of the invention, the compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof may be administered in association with an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and/or a bile acid binder thereby avoiding a possible risk of excess of bile acids in colon caused by the inhibition of the ileal bile acid transport system. An excess of bile acids in the visceral contents may cause diarrhoea. Thus, the present invention also provides a treatment of a possible side effect such as diarrhoea in patients during therapy comprising the compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

An HMG CoA-reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof will by its action decrease the endogenous cholesterol available for the bile acid synthesis and have an additive effect in combination with the compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof on lipid lowering.

Suitable bile acid binders for such a combination therapy are resins, such as cholestyramine and cholestipol. One advantage is that the dose of bile acid binder might be kept lower than the therapeutic dose for treatment of cholesterolaemia in single treatment comprising solely a bile acid binder. By a low dose of bile acid binder any possible side effects caused by poor tolerance of the patient to the therapeutic dose could also be avoided.

Therefore in an additional feature of the invention, there is provided a method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore in an additional feature of the invention, there is provided a method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with a bile acid binder.

Therefore in an additional feature of the invention, there is provided a method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in simultaneous, sequential or separate administration with a bile acid binder.

Therefore in an additional feature of the invention, there is provided a method of treating hyperlipidemic conditions in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore in an additional feature of the invention, there is provided a method of treating hyperlipidemic conditions in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of a bile acid binder.

Therefore in an additional feature of the invention, there is provided a method of treating hyperlipidemic conditions in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in simultaneous, sequential or separate administration with a bile acid binder.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder, in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the present invention there is provided a kit comprising a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

According to a further aspect of the present invention there is provided a kit comprising a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder.

According to a further aspect of the present invention there is provided a kit

5 comprising a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof and a bile acid binder.

According to a further aspect of the present invention there is provided a kit

10 comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;
- b) an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof; in a second unit dosage form; and

15 c) container means for containing said first and second dosage forms.

According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;
- b) a bile acid binder; in a second unit dosage form; and
- c) container means for containing said first and second dosage forms.

According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;
- b) an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof; in a second unit dosage form;
- c) a bile acid binder; in a third unit dosage form; and
- d) container means for containing said first, second and third dosage forms.

30 According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a second unit dosage form; and
- c) container means for containing said first and second dosage forms.

According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) a bile acid binder, in a second unit dosage form; and
- c) container means for containing said first and second dosage forms.

According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a second unit dosage form; and
- c) a bile acid binder; in a third unit dosage form; and
- d) container means for containing said first, second and third dosage forms.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder, in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a

prodrug thereof, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder, in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

5 According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in the manufacture of a medicament for use in the treatment of hyperlipidaemic conditions in a warm-blooded animal, such as man.

10 According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, a bile acid binder, in the manufacture of a medicament for use in the treatment of hyperlipidaemic conditions in a warm-blooded animal, such as man.

According to another feature of the invention there is provided the use of a compound
15 of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder, in the manufacture of a medicament for use in the treatment of hyperlipidaemic conditions in a warm-blooded animal, such as man.

20 According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, with the simultaneous, sequential or separate administration of an effective amount of an HMG Co-A
25 reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of a compound of the formula
30 (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, with the simultaneous, sequential or separate administration of an effective amount of a bile acid

binder, optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of a compound of the formula 5 (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, with the simultaneous, sequential or separate administration of an effective amount of an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable excipient, with the 10 simultaneous, sequential or separate administration of an effective amount of a bile acid binder, optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment.

According to an additional further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of a compound 15 of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, with the simultaneous, sequential or separate administration one or more of the following agents selected from:

- a CETP (cholesteryl ester transfer protein) inhibitor, for example those referenced and 20 described in WO 00/38725 page 7 line 22 - page 10, line 17 which are incorporated herein by reference;
- a cholesterol absorption antagonist for example azetidinones such as SCH 58235 and those described in US 5,767,115 which are incorporated herein by reference;
- a MTP (microsomal transfer protein) inhibitor for example those described in Science, 25 282, 751-54, 1998 which are incorporated herein by reference;
- a fibric acid derivative; for example clofibrate, gemfibrozil, fenofibrate, ciprofibrate and bezafibrate;
- a nicotinic acid derivative, for example, nicotinic acid (niacin), acipimox and nericitrol;
- 30 ➤ a phytosterol compound for example stanols;
- probucol;
- an anti-obesity compound for example orlistat (EP 129,748) and sibutramine (GB 2,184,122 and US 4,929,629);

- an antihypertensive compound for example an angiotensin converting enzyme (ACE) inhibitor, an angiotensin II receptor antagonist, an adrenergic blocker, an alpha adrenergic blocker, a beta adrenergic blocker, a mixed alpha/beta adrenergic blocker, an adrenergic stimulant, calcium channel blocker, a diuretic or a vasodilator;
- 5 ➤ insulin;
- sulphonylureas including glibenclamide, tolbutamide;
- metformin; and/or
- acarbose;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof,

10 optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment.

Particular ACE inhibitors or pharmaceutically acceptable salts, solvates, solvate of such salts or a prodrugs thereof, including active metabolites, which can be used in combination with a compound of formula (I) include but are not limited to, the following

15 compounds: alacepril, alatriopril, altiopril calcium, ancovenin, benazepril, benazepril hydrochloride, benazeprilat, benzoylcaptopril, captopril, captopril-cysteine, captopril-glutathione, ceranapril, ceranopril, ceronapril, cilazapril, cilazaprilat, delapril, delapril-diacid, enalapril, enalaprilat, enapril, epicaptopril, foroxymithine, fosfenopril, fosenopril, fosenopril sodium, fosinopril, fosinopril sodium, fosinoprilat, fosinoprilic acid, glycopril, hemorphin-4,

20 idrapril, imidapril, indolapril, indolaprilat, libenzapril, lisinopril, lyciumin A, lyciumin B, mixanpril, moexipril, moexiprilat, moveltipril, muracein A, muracein B, muracein C, pentopril, perindopril, perindoprilat, pivalopril, pivopril, quinapril, quinapril hydrochloride, quinaprilat, ramipril, ramiprilat, spirapril, spirapril hydrochloride, spiraprilat, spiropril, spiropril hydrochloride, temocapril, temocapril hydrochloride, teprotide, trandolapril,

25 trandolaprilat, utibapril, zabicipril, zabiciprilat, zofenopril and zofenoprilat. Preferred ACE inhibitors for use in the present invention are ramipril, ramiprilat, lisinopril, enalapril and enalaprilat. More preferred ACE inhibitors for uses in the present invention are ramipril and ramiprilat.

Preferred angiotensin II antagonists, pharmaceutically acceptable salts, solvates,

30 solvate of such salts or a prodrugs thereof for use in combination with a compound of formula (I) include, but are not limited to, compounds: candesartan, candesartan cilexetil, losartan, valsartan, irbesartan, tasosartan, telmisartan and eprosartan. Particularly preferred angiotensin

II antagonists or pharmaceutically acceptable derivatives thereof for use in the present invention are candesartan and candesartan cilexetil.

In another aspect of the invention, the compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, may be administered in association with a PPAR alpha and/or gamma agonist, or pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof. Suitable PPAR alpha and/or gamma agonists, pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof are well known in the art. These include the compounds described in WO 01/12187, WO 01/12612, WO 99/62870, WO 99/62872, WO 99/62871, WO 98/57941, WO 01/40170, J Med Chem, 1996, 39, 665, Expert Opinion on Therapeutic Patents, 10 (5), 623-634 (in particular the compounds described in the patent applications listed on page 634) and J Med Chem, 2000, 43, 527 which are all incorporated herein by reference. Particularly a PPAR alpha and/or gamma agonist refers to WY-14643, clofibrate, fenofibrate, bezafibrate, GW 9578, troglitazone, pioglitazone, rosiglitazone, eglitazone, proglitazone, BRL-49634, KRP-297, JTT-501, SB 213068, GW 1929, GW 7845, GW 0207, L-796449, L-165041 and GW 2433. Particularly a PPAR alpha and/or gamma agonist refers to (S)-2-ethoxy-3-[4-(2-{4-methanesulphonyloxyphenyl}ethoxy)phenyl] propanoic acid and pharmaceutically acceptable salts thereof.

Therefore in an additional feature of the invention, there is provided a method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore in an additional feature of the invention, there is provided a method of treating hyperlipidemic conditions in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in simultaneous, sequential or separate administration with an effective amount of a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug

5 thereof, in association with a pharmaceutically acceptable diluent or carrier.

According to a further aspect of the present invention there is provided a kit comprising a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

10 According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;
- b) a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof; in a second unit dosage form; and
- c) container means for containing said first and second dosage forms.

According to a further aspect of the present invention there is provided a kit comprising:

- a) a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, together with a pharmaceutically acceptable diluent or carrier, in a first unit dosage form;
- b) a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a second unit dosage form; and
- c) container means for containing said first and second dosage forms.

25 According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as

30 man.

According to another feature of the invention there is provided the use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt,

solvate, solvate of such a salt or a prodrug thereof, in the manufacture of a medicament for use in the treatment of hyperlipidaemic conditions in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of a compound of the formula

5 (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, with the simultaneous, sequential or separate administration of an effective amount of a PPAR alpha and/or gamma agonist, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier to
10 a warm-blooded animal, such as man in need of such therapeutic treatment.

In addition to their use in therapeutic medicine, the compounds of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, are also useful as pharmacological tools in the development and standardisation of *in vitro* and *in vivo* test systems for the evaluation of the effects of inhibitors of IBAT in laboratory animals such
15 as cats, dogs, rabbits, monkeys, rats and mice, as part of the search for new therapeutic agents.

Many of the intermediates described herein are novel and are thus provided as a further feature of the invention. For example compounds of formula (VIIIa), (VIIIb), (IXa),
(IXb), (XIIIa) and (XIIIb) show IBAT inhibitory activity when tested in the above referenced *in vitro* test assay and are thus claimed as a further feature of the invention.

20 Thus in a further feature of the invention, there is provided a compound of formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Therefore according to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of formula (VIIIa), (VIIIb),
25 (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore in association with a pharmaceutically-acceptable diluent or carrier.

According to an additional aspect of the present invention there is provided a compound of the formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a
30 pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore for use in a method of prophylactic or therapeutic treatment of a warm-blooded animal, such as man.

Thus according to this aspect of the invention there is provided a compound of the formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as defined hereinbefore for use as a medicament.

5 According to another feature of the invention there is provided the use of a compound of the formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof as defined hereinbefore in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

10 According to another feature of the invention there is provided the use of a compound of the formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof as defined hereinbefore in the manufacture of a medicament for use in the treatment of hyperlipidaemic conditions in a warm-blooded animal, such as man.

15 According to a further feature of this aspect of the invention there is provided a method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

20 According to a further feature of this aspect of the invention there is provided a method of treating hyperlipidemic conditions in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

25 In the above other pharmaceutical composition, process, method, use and medicament manufacture features, the alternative and preferred embodiments of the compounds of the invention described herein also apply.

Examples

30 The invention will now be illustrated in the following non limiting Examples, in which standard techniques known to the skilled chemist and techniques analogous to those described in these Examples may be used where appropriate, and in which, unless otherwise stated:

(i) evaporation were carried out by rotary evaporation in vacuo and work up procedures were carried out after removal of residual solids such as drying agents by filtration;

(ii) all reactions were carried out under an inert atmosphere at ambient temperature, typically in the range 18-25°C, with solvents of HPLC grade under anhydrous conditions, unless

5 otherwise stated;

(iii) column chromatography (by the flash procedure) was performed on Silica gel 40-63 µm (Merck);

(iv) yields are given for illustration only and are not necessarily the maximum attainable;

(v) the structures of the end products of the formula (I) were generally confirmed by nuclear

10 (generally proton) magnetic resonance (NMR) and mass spectral techniques; magnetic resonance chemical shift values were measured in deuterated CDCl₃ (unless otherwise stated) on the delta scale (ppm downfield from tetramethylsilane); proton data is quoted unless otherwise stated; spectra were recorded on a Varian Mercury-300 MHz, Varian Unity plus-400 MHz, Varian Unity plus-600 MHz or on Varian Inova-500 MHz spectrometer unless

15 otherwise stated data was recorded at 400MHz; and peak multiplicities are shown as follows: s, singlet; d, doublet; dd, double doublet; t, triplet; tt, triple triplet; q, quartet; tq, triple quartet; m, multiplet; br, broad; ABq, AB quartet; ABD, AB doublet, ABdd, AB doublet of doublets; dABq, doublet of AB quartets; LCMS were recorded on a Waters ZMD, LC column xTerra

20 MS C₈(Waters), detection with a HP 1100 MS-detector diode array equipped; mass spectra (MS) (loop) were recorded on VG Platform II (Fisons Instruments) with a HP-1100 MS-detector diode array equipped; unless otherwise stated the mass ion quoted is (MH⁺); unless further details are specified in the text, analytical high performance liquid chromatography (HPLC) was performed on Prep LC 2000 (Waters), Cromasil C₈, 7 µm, (Akzo Nobel); MeCN and de-ionised water 10 mM ammonium acetate as mobile phases, with

25 suitable composition;

(vii) intermediates were not generally fully characterised and purity was assessed by thin layer chromatography (TLC), HPLC, infra-red (IR), MS or NMR analysis;

(viii) where solutions were dried sodium sulphate was the drying agent;

(ix) where an "ISOLUTE" column is referred to, this means a column containing 2 g of silica,

30 the silica being contained in a 6 ml disposable syringe and supported by a porous disc of 54Å pore size, obtained from International Sorbent Technology under the name "ISOLUTE"; "ISOLUTE" is a registered trade mark;

(x) the following abbreviations may be used hereinbefore or hereinafter:-

DCM	dichloromethane;
DMF	<i>N,N</i> -dimethylformamide;
TBTU	o-Benzotriazol-1-yl- <i>N,N,N',N'</i> -tetramethyluronium tetrafluoroborate;
EtOAc	EtOAC;
5 MeCN	acetonitrile;
TFA	trifluoroacetic acid;
IPA	isopropanol;
DIPEA	di-isopropylethylamine; and
THF	tetrahydrofuran.

10

Example 1

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-

15 methoxycarbonylmethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 29; 300 mg, 0.46 mmol) was dissolved in methanol (5 ml). NaOH (100 mg in 0.2 ml water) was added to the solution and the mixture was stirred at room temperature for 1 hour. Acetic acid (0.3 ml) was added. The solvent was evaporated under reduced pressure and the residue was extracted with DCM/water. The DCM layer was separated, dried and evaporated under 20 reduced pressure to give the title compound 270 mg (92%). NMR, 500 MHz 0.7-0.8 (m, 6H), 1.0-1.6 (m, 12H), 2.1 (s, 3H) 3.2 (brs, 2H), 3.6-3.8 (m, 2H), 4.6 (s, 2H), 5.6 (d, 1H), 6.6 (s, 1H), 6.9-7.5 (m, 11H), 7.8 (d, 1H).

Examples 2-9

25 The following compounds were synthesised by the procedure of Example 1 using the appropriate starting material.

Ex	Compound	NMR	SM
2		(300 MHz, CD ₃ OD) 0.9-0.95 (m, 6H), 1.05-1.3 (m, 8H), 1.4-1.6 (m, 4H), 2.2 (s, 3H), 3.25 (s, 2H), 3.75 (brs, 2H), 4.65 (dd, 2H), 5.2 (s, 1H), 6.7-7.3 (m, 10H), 7.4 (s, 1H)	Meth 30

3		(300 MHz, CD ₃ OD) 0.75-0.85 (m, 6H), 1.0-1.6 (m, 12H), 2.2 (s, 3H), 3.75 (brs, 2H), 3.25 (s, 2H), 4.6-4.7 (m, 2H), 5.7 (s, 1H), 6.7 (s, 1H), 6.9-7.3 (m, 8H), 7.4 (s, 1H)	Meth 31
4		(300 MHz, CD ₃ OD) 0.75-0.9 (m, 6H), 1.0-1.6 (m, 12H), 2.2 (s, 3H), 3.25 (s, 2H), 3.75 (brs, 2H), 4.6-4.8 (m, 2H), 5.45 (s, 1H), 6.7 (s, 1H), 6.95-7.3 (m, 5H), 7.4 (s, 1H), 7.6 (s, 4H)	Meth 32
5		(500 MHz) 0.7-0.8 (m, 6H), 1.0-1.6 (m, 12H), 3.2 (brs, 2H), 3.6 (brs, 2H), 4.48 (m, 2H), 5.0 (s, 1H), 6.5 (d, 1H), 6.7-7.4 (m, 10H), 7.9 (s, 1H)	Meth 39
6		(DMSO-d ₆) 0.7-0.8 (m, 6H), 0.9-1.6 (m, 12H), 3.2 (brs, 2H), 3.7 (brs, 2H), 4.6-4.8 (m, 3H), 6.6 (d, 2H), 6.9-7.3 (m, 8H), 7.4 (s, 1H), 8.3 (d, 1H)	Meth 40
7		M/z = 768.9	Meth 67

¹		(300 MHz, CD ₃ OD) 0.75-0.9 (m, 6H), 1.0-1.25 (m, 4H), 1.4-1.6 (m, 4H), 2.15 (s, 3H), 3.1-3.3 (m, 4H), 3.5-3.8 (m, 5H), 4.75 (ABq, 2H), 5.45 (s, 2H), 6.75 (s, 1H), 6.95-7.5 (11H); m/z 711.3	Meth 42
9		(500 MHz, DMSO-d ₆) 0.7-0.8 (m, 6H), 0.9-1.6 (m, 12H), 2.2 (s, 3H) 3.2-3.8 (m, 8H), 4.8 (ABq, 2H), 5.6 (d, 1H), 6.7 (s, 1H), 6.8-7.5 (m, 11H), 7.8 (brs, 1H), 8.6 (d, 1H), 8.8 (t, 1H)	Meth 69

¹ Ethanol instead of methanol, purified by preparative HPLC using MeCN and ammonium acetate buffer (55:45) as eluent

Example 10

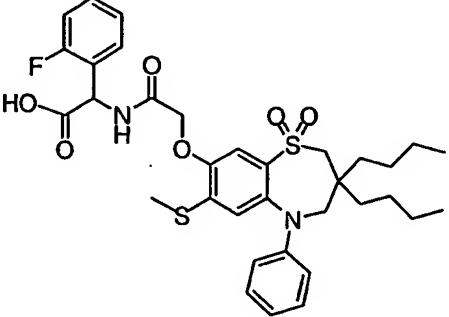
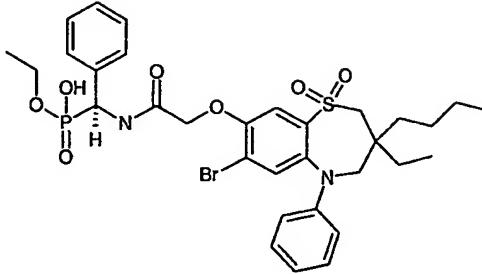
5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-{1-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoyl]ethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-{1-[N-((R)-1'-phenyl-1'-methoxycarbonylmethyl)carbamoyl]ethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 33; 103 mg, 0.15 mmol) was dissolved in a mixture of THF and H₂O (2:1, 3 ml). LiOH (7 mg, 0.3 mmol) was added and the mixture was stirred for 7 hours at ambient temperature. Most of the solvent was removed under reduced pressure and the crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (45:55) as eluent to give the title compound 97 mg (96 %). NMR (DMSO-d₆) 0.60-0.80 (m, 6H), 0.90-1.60 (m, 11H), 3.15-3.45 (m, 2H), 3.50-3.90 (m, 2H), 4.95-5.25 (m, 2H), 6.85-7.55 (m, 12H), 8.55-8.95 (m, 1H).

Examples 11-16

The following compounds were synthesised by the procedure of Example 10 using the appropriate starting material.

Ex	Compound	NMR	SM
11		((CD ₃) ₂ CO) 0.70-0.90 (m, 6H), 0.95-1.35 (m, 4H), 1.40-1.75 (m, 4H), 3.15-3.35 (m, 2H), 3.80 (brs, 2H), 5.40 (d, 1H), 5.90-6.15 (2s, 1H), 6.95-7.75 (m, 18H)	Meth 34
12		(CD ₃ OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.55 (m, 4H), 3.20 (s, 2H), 3.60 (s, 3H), 3.75 (brs, 2H), 4.60 (ABq, 2H), 5.40 (s, 1H), 6.50 (s, 1H), 6.95-7.45 (m, 10H), 7.55 (s, 1H)	Meth 35
13		(CD ₃ OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.55 (m, 4H), 3.20 (s, 2H), 3.55 (s, 3H), 3.75 (brs, 2H), 3.90 (ABq, 2H), 4.60 (ABq, 2H), 5.60 (s, 1H), 6.50 (s, 1H), 6.95-7.45 (m, 10H), 7.55 (s, 1H)	Meth 36
14		(CD ₃ OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.60 (m, 4H), 3.20 (s, 2H), 3.60 (s, 3H), 3.75 (brs, 2H), 4.55 (ABq, 2H), 5.55 (s, 1H), 6.50 (s, 1H), 6.95-7.45 (m, 9H), 7.50 (s, 1H)	Meth 37

15		(CD ₃ OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.60 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.75 (brs, 2H), 4.65 (ABq, 2H), 5.60 (s, 1H), 6.70 (s, 1H), 6.90-7.45 (m, 10H)	Meth 38
16		(CD ₃ OD) 7.55-7.41 (3H, m), 7.35-7.20 (5H, m), 7.15-7.08 (3H, m), 7.04-6.98 (1H, m), 5.48-5.32 (1H, m), 4.80-4.60 (2H, m), 4.00-3.56 (4H, m), 3.27-3.22 (2H, m), 1.61-1.00 (11H, m), 0.83-0.74 (6H, m)	Meth 70

Example 17**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((S)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

5. 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((S)-1'-phenyl-1'-methoxycarbonylmethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 46; 60 mg, 0.091 mmol) was dissolved in THF (1 ml) and added to a solution of lithium hydroxide monohydrate (12.6 mg, 0.29 mmol) in water (1 ml). The mixture was stirred occasionally for 30 minutes. 2M HCl solution (0.3 ml) was added and the water layer was 10 extracted with DCM. The organic layer was washed once with brine, dried, filtered and evaporated at reduced pressure to give the title compound 48 mg (82%). NMR (CD₃OD) 0.73-0.84 (m, 6H), 1.0-1.6 (m, 8H), 3.27 (brs, 2H), 3.60-3.90 (m, 2H), 4.71 (ABq, 2H), 5.47-5.55 (m, 1H), 7.02 (brt, 1H), 7.08-7.17 (m, 3H), 7.25-7.46 (m, 7H), 7.52 (s, 1H), 8.43 (d, NH); m/z 643.5.

15

Examples 18-21

The following compounds were synthesised by the procedure of Example 17 using the appropriate starting material.

Ex	Compound	NMR or m/z	SM
18 1		M/z 670 (M+NH4 ⁺)	Meth 43
19 2		(CD ₃ OD) 0.70-0.90 (m, 6H), 1.0-1.32 (m, 4H), 1.32-1.70 (m, 4H), 2.15 (s, 3H), 2.85 (brs, 3H), 3.23 (brs, 2H), 3.53-3.93 (m, 2H), 4.99 (ABq, 2H), 6.27 (s, 1H), 6.71 (s, 1H), 6.94 (t, 1H), 7.07 (d, 2H), 7.25 (t, 2H), 7.3-7.47 (m, 6H); m/z 625.3	Meth 62
20		(CD ₃ OD) 0.75-0.84 (m, 6H), 1.0-1.29 (m, 4H), 1.36-1.65 (m, 4H), 2.15 (s, 3H), 2.82-2.97 (m, 2H), 3.22 (brs, 2H), 3.6-3.85 (m, 2H), 4.66 (ABq, 2H), 5.43 (t, 1H), 6.71 (s, 1H), 6.96 (t, 1H), 7.09 (d, 2H), 7.2-7.38 (m, 7H), 7.40 (s, 1H); m/z 625.4	Meth 112
21		(600 MHz, CD ₃ OD) 0.77-0.88 (m, 6H), 1.0-1.32 (m, 4H), 1.39-1.70 (m, 4H), 2.16 (s, 3H), 2.88 (brs, 3H), 3.25 (brs, 2H), 3.52-3.93 (m, 2H), 5.03 (ABq, 2H), 6.28 (s, 1H), 6.73 (s, 1H), 6.96 (t, 1H), 7.09 (brd, 2H), 7.27 (t, 2H), 7.32-7.46 (m, 6H)	Meth 79

2.2 equivalents of LiOH in THF/water (4/1)

² Purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent

Example 22

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

The title compound was synthesised from 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-methoxycarbonylmethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 61) by the procedure of Example 17, except that the
10 water layer was extracted with EtOAc. The product was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent. NMR 0.75-0.83 (m, 6H), 1.0-1.25 (m, 4H), 1.32-1.52 (m, 3H), 1.55-1.70 (m, 1H), 3.20 (ABq, 2H), 3.65-3.83 (m, 2H), 4.62 (ABq, 2H), 5.68 (d, 1H), 7.04-7.15 (m, 4H), 7.3-7.5 (m, 8H), 7.87 (brd, 1H); m/z 643.1.

15

Example 23

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-(N-[(S)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine ammonium salt

20 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((S)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 17; 48 mg, 0.075 mmol) and 2-aminoethanesulphonic acid (17 mg, 0.14 mmol) was dissolved in DMF (2 ml) and DIPEA (0.052 ml, 0.30 mmol). The mixture was stirred for 15 min at 60°C. TBTU (31 mg, 0.097 mmol) was added and the mixture was stirred for 2 hours at 60°C. The
25 solvent was evaporated at reduced pressure. The residue was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent. Lyophilisation gave the title compound 4 mg (7%). NMR (CD₃OD) 0.75-0.83 (m, 6H), 0.95-1.65 (m, 8H), 2.85-3.0 (m, 2H), 3.27 (brs, 2H), 3.5-3.9 (m, 4H), 4.72 (ABq, 2H), 5.48 (s, 1H), 7.02 (brt, 1H), 7.09-7.15 (m, 3H), 7.25-7.52 (m, 8H); m/z 750.3.

30

Examples 24-37

The following compounds were prepared by the same procedure. The acid (1 equiv) was dissolved in THF (1 ml) and added to a solution of lithium hydroxide monohydrate (12.6 mg, 2.9-6.6 equiv) in water (1 ml). The mixture was stirred occasionally and after 1.5-6 hours 5 the deprotection was completed (according to LC-MS). 2M HCl-solution (0.3 ml) was added.

Examples 24-33

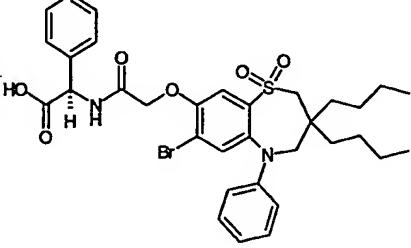
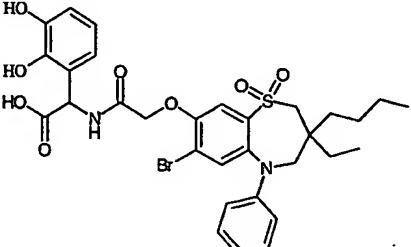
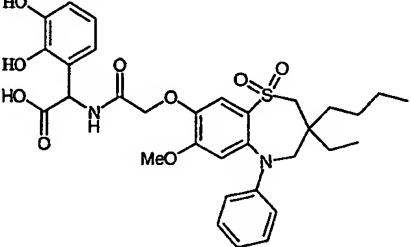
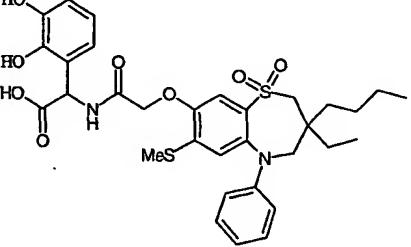
The reaction mixture was put on a syringe filled with hydramatrix®. The product was eluted with DCM. The DCM was dried, filtered and evaporated at reduced pressure. The product was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient 10 (5/95 to 100/0) as eluent.

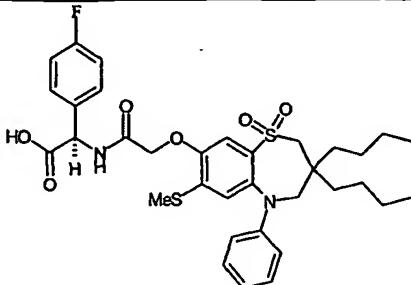
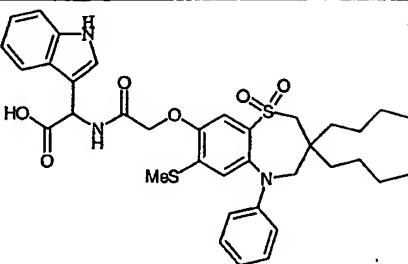
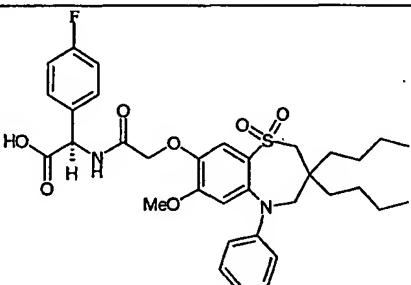
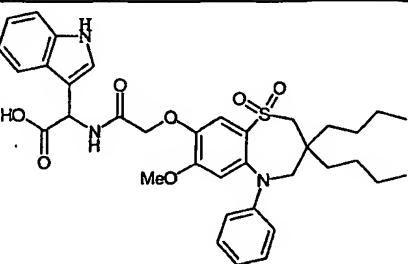
Examples 34-37

The water layer was extracted two times with DCM. The organic layer was dried, filtered and evaporated at reduced pressure.

Ex	Compound	NMR (CD_3OD)	m/z	SM
24		0.75-0.84 (m, 6H), 1.0-1.25 (m, 4H), 1.37-1.65 (m, 4H), 3.20 (brs, 2H), 3.55-3.90 (m, 5H), 4.58 (ABq, 2H), 5.33 (s, 1H), 6.51 (s, 1H), 6.97 (brt, 1H), 7.12 (brd, 2H), 7.2-7.33 (m, 5H), 7.41 (brd, 2H), 7.54 (s, 1H)	595.4 47	Meth 47
25		0.73-0.85 (m, 6H), 1.0-1.3 (m, 4H), 1.35-1.65 (m, 4H), 2.17 (s, 3H), 3.23 (brs, 2H), 3.55-3.90 (m, 2H), 4.71 (ABq, 2H), 5.49-5.52 (m, 1H), 6.73 (s, 1H), 6.96 (brt, 1H), 7.10 (brd, 2H), 7.23-7.45 (m, 8H), 8.36 (brd, NH)	611.2 48	Meth 48

26		0.74-0.84 (m, 6H), 1.0-1.3 (m, 8H), 1.37-1.54 (m, 4H), 3.28 (brs, 2H), 3.65-3.85 (m, 2H), 4.72 (ABq, 2H), 5.49-5.52 (m, 1H), 7.04 (brt, 1H), 7.09-7.18 (m, 3H), 7.28-7.46 (m, 7H), 7.52 (s, 1H), 8.45 (brd, NH)	671.2	Meth 49
27		0.74-0.84 (m, 6H), 1.0-1.3 (m, 4H), 1.35-1.65 (m, 4H), 3.21 (brs, 2H), 3.59 (s, 3H), 3.62-3.90 (m, 2H), 4.62 (ABq, 2H), 5.49 (s, 1H), 6.50 (s, 1H), 6.98 (brt, 1H), 7.12 (brd, 2H), 7.24-7.43 (m, 7H), 7.54 (s, 1H)	595.3	Meth 50
28		0.74-0.85 (m, 6H), 0.85-1.65 (m, 14H), 3.21 (brs, 2H), 3.6-3.9 (m, 2H), 4.25-4.36 (m, 1H), 4.53-4.66 (m, 2H), 5.49 (s, 1H), 6.47 (s, 1H), 6.91-7.0 (m, 1H), 7.04-7.16 (m, 2H), 7.22-7.46 (m, 7H), 7.51 (s, 1H)	623.3	Meth 51
29		0.73-0.85 (m, 6H), 0.85-1.65 (m, 8H), 3.24 (brs, 3H), 3.34 (brs, 2H), 3.6-3.95 (m, 2H), 4.8-4.95 (m, 2H), 5.52 (s, 1H), 7.06 (brt, 1H), 7.17 (brd, 2H), 7.27-7.40 (m, 5H), 7.40-7.50 (m, 3H), 7.69 (s, 1H)	643.3	Meth 52

30		0.74-0.84 (m, 6H), 0.85-1.55 (m, 12H), 3.24-3.33 (m, 2H), 3.65-3.85 (m, 2H), 4.65-4.78 (m, 2H), 5.50 (brs, 1H), 6.99-7.2 (m, 4H), 7.25-7.48 (m, 7H), 7.51 (s, 1H)	671.2	Meth 53
31		0.72-0.84 (m, 6H), 0.85-1.65 (m, 8H), 3.27 (brs, 2H), 3.54-3.9 (m, 2H), 4.70 (ABq, 2H), 5.70 (s, 1H), 6.63-6.69 (m, 1H), 6.71-6.77 (m, 2H), 7.02 (brt, 1H), 7.08-7.17 (m, 3H), 7.30 (brt, 2H), 7.52 (s, 1H)	675.4	Meth 54
32		0.72-0.84 (m, 6H), 0.98-1.67 (m, 8H), 3.21 (brs, 2H), 3.54-3.9 (m, 5H), 4.62 (ABq, 2H), 5.57 (s, 1H), 6.51 (s, 1H), 6.59-6.73 (m, 3H), 6.97 (brt, 1H), 7.12 (brd, 2H), 7.28 (brt, 2H), 7.56 (s, 1H)	627.5	Meth 55
33		0.73-0.86 (m, 6H), 1.0-1.68 (m, 8H), 2.19 (s, 3H), 3.24 (brs, 2H), 3.55-3.9 (m, 2H), 4.71 (ABq, 2H), 5.53 (s, 1H), 6.60-6.73 (m, 3H), 6.75 (s, 1H), 6.96 (brt, 1H), 7.10 (brd, 2H), 7.27 (brt, 2H), 7.44 (s, 1H)	643.4	Meth 56

34		0.74-0.86 (m, 6H), 1.0-1.3 (m, 8H), 1.35-1.57 (m, 4H), 2.19 (s, 3H), 3.23 (brs, 2H), 3.62-3.85 (m, 2H), 4.65 (ABq, 2H), 5.28 (s, 1H), 6.72 (s, 1H), 6.94-7.05 (m, 3H), 7.12 (brd, 2H), 7.28 (brt, 2H), 7.39 (s, 1H), 7.43 (dd, 2H)	657.3	Meth 57
35		0.75-0.86 (m, 6H), 1.0-1.3 (m, 8H), 1.35-1.55 (m, 4H), 2.01 (s, 3H), 3.11-3.26 (ABq, 2H), 3.6-3.8 (m, 2H), 4.58 (d, 1H), 4.70 (d, 1H), 5.64 (s, 1H), 6.62 (s, 1H), 6.91-7.0 (m, 2H), 7.01-7.12 (m, 3H), 7.23-7.33 (m, 4H), 7.37 (s, 1H), 7.69 (brd, 1H)	678.4	Meth 58
36		0.76-0.84 (m, 6H), 1.0-1.3 (m, 8H), 1.36-1.53 (m, 4H), 3.21 (brs, 2H), 3.64 (s, 3H), 3.67-3.87 (m, 2H), 4.57 (ABq, 2H), 5.31 (s, 1H), 6.50 (s, 1H), 6.95-7.06 (m, 3H), 7.14 (brd, 2H), 7.28 (brt, 2H), 7.38-7.46 (m, 2H), 7.51 (s, 1H)	641.4	Meth 59
37		0.75-0.87 (m, 6H), 1.0-1.3 (m, 8H), 1.34-1.53 (m, 4H), 3.18 (ABq, 2H), 3.27 (s, 3H), 3.65-3.85 (m, 2H), 4.52 (d, 1H), 4.65 (d, 1H), 5.66 (s, 1H), 6.30 (s, 1H), 6.90-7.02 (m, 2H), 7.03-7.16 (m, 3H), 7.23-7.34 (m, 4H), 7.50 (s, 1H), 7.59 (brd, 1H)	662.4	Meth 60

Example 38

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 22; 50 mg, 0.078 mmol) was dissolved in DMF (1.5 ml). Sodium methanethiolate (20 mg, 0.29 mmol) was added and the mixture was stirred for 1.5 hours at 50°C. Acetic acid (40 mg) was added and the solvent was evaporated under reduced pressure. The residue was purified by preparative HPLC using MeCN/ammonium acetate buffer (45:55) as eluent to give the title compound 29 mg (61%). NMR (DMSO-d₆): 0.7-0.8 (m, 6H), 0.9-1.6 (m, 8H), 2.2 (s, 3H), 3.1-3.7 (m, 4H), 4.6-4.8 (m, 3H), 6.7 (s, 1H), 6.8-7.4 (m, 11H), 8.3 (d, 1H).

Example 39

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-ethylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 22; 50 mg, 0.078 mmol), ethanethiol (99 mg, 1.59 mmol) and caesium carbonate (253 mg, 0.78 mmol) were added to DMF (5 ml) and the mixture was stirred for 30 h at 44°C. The solvent was filtered, evaporated under reduced pressure. The residue was purified by preparative HPLC using MeCN/ammonium acetate buffer (45:55) as eluent. The residue was purified by column chromatography using DCM : methanol (100:15) to give the title compound 15 mg (31%). NMR (300 MHz, CD₃OD) 0.7-0.85 (m, 6H), 1.0-1.6 (m, 11H), 2.65 (q, 2H), 3.2 (s, 2H), 3.7 (brs, 2H), 4.6 (q, 2H), 5.3 (s, 1H), 6.75 (s, 1H), 6.9-7.5 (m, 11H).

25

Example 40

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-(2-hydroxyethylthio)-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 22; 50 mg, 0.078 mmol), 2-mercaptopropanol (281 mg, 3.59 mmol) and caesium carbonate (228 mg, 0.7 mmol) were added to DMF (5 ml) and the mixture was stirred for 9 hours at 70°C. The solvent was evaporated under reduced pressure. The residue was purified by preparative

HPLC using MeCN/ammonium acetate buffer (45:55) as eluent. The collected fractions were lyophilised to give the title compound 20 mg (40%). NMR (300 MHz, CD₃OD) 0.75-0.85 (m, 6H), 1.0-1.6 (m, 8H), 2.9 (t, 2H), 3.2 (s, 2H), 3.55 (t, 2H), 3.7 (brs, 2H), 4.65 (q, 2H), 5.3 (s, 1H), 6.9 (s, 1H), 6.95-7.5 (m, 11H).

5

Example 41

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-(2-N,N'-dimethylaminoethylthio)-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 22; 50 mg, 0.078 mmol), dimethylaminoethanethiol hydrochloride (99 mg, 0.94 mmol), potassium carbonate (129 mg, 0.94 mmol), DIPEA (100 mg, 0.77 mmol) and sodium borohydride (35 mg, 0.93 mmol) were added to DMF (10 ml) and the mixture was stirred for 24 hours at 85°C. The solvent was filtered and evaporated under reduced pressure. The residue was purified twice by preparative HPLC using MeCN/ammonium acetate buffer (40:60) as eluent. The collected fractions were lyophilised to give the title compound 15 mg (30%). NMR (300 MHz, CD₃OD) 0.75-0.85 (m, 6H), 1.0-1.65 (m, 8H), 2.65 (s, 6H), 3.05 (t, 2H), 3.2 (t, 2H), 3.3 (s, 2H), 3.75 (brs, 2H), 4.75 (s, 2H), 5.2 (s, 1H), 6.95-7.4 (m, 11), 7.5 (s, 1H).

20 **Example 42**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-isopropylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 22; 50 mg, 0.078 mmol), 2-propanethiol (126 mg, 1.65 mmol), caesium carbonate (152 mg, 0.47 mmol), sodium borohydride (25 mg, 0.66 mmol) were added to DMF (5 ml) and the mixture was stirred for 5 min at 100°C. The solvent was evaporated under reduced pressure. The residue was purified by preparative HPLC using MeCN/ammonium acetate buffer (45:55) as eluent. The collected fractions were lyophilised to give the title compound 15 mg (30%).

30 NMR (300 MHz, DMSO-d₆) 0.7-0.85 (m, 6H), 0.95-1.65 (m, 14H), 3.3 (s, 2H), 3.7 (brs, 2H), 4.75 (dd, 2H), 5.05 (brs, 1H), 6.75-7.4 (m, 12H), 8.5 (brs, 1H).

Example 43

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(carboxymethyl)carbamoylmethyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(*t*-butoxycarbonylmethyl)carbamoylmethyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 63; 120 mg, 0.17 mmol) was dissolved in DCM (2 ml). TFA (0.7 ml) was added and the mixture was stirred at room temperature for 3 h. The reaction mixture was evaporated under reduced pressure. The residue was purified by preparative HPLC using
10 MeCN/ammonium acetate buffer (50:50) as eluent to give the title compound 95 mg (85%).
NMR (300 MHz, DMSO-d₆) 0.7-0.8 (m, 6H), 0.9-1.6 (m, 12H), 2.2 (s, 3H) 3.2-3.3 (m, 2H), 3.5-3.8 (m, 4H), 4.8 (ABq, 2H), 5.6 (d; 1H), 6.7 (s, 1H), 6.8-7.5 (m, 11H), 8.5-8.7 (m, 2H).

Examples 44-49

15 The following compounds were synthesised by the procedure of Example 43 using the appropriate starting material.

Ex	Compound	NMR or m/z	SM
44		(300 MHz) 0.7-0.9 (m, 6H), 1.0-1.7 (m, 8H), 3.2 (m, 2H), 3.75 (brs, 2H), 3.9-4.0 (m, 1H), 4.15-4.25 (m, 1H), 4.5-4.7 (m, 2H), 5.75-5.9 (m, 1H), 7.05-7.2 (m, 4H), 7.25-7.4 (m, 5H), 7.45-7.55 (m, 3H), 8.2 (d, 1H)	Meth 64
45		(CD ₃ OD) 0.70-0.90 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.55 (m, 4H), 3.20 (s, 2H), 3.55 (s, 3H), 3.75 (brs, 2H), 3.80-4.00 (m, 2H), 4.40-4.70 (m, 3H), 5.65 (s, 1H), 6.50 (s, 1H), 6.95-7.50 (m, 10H), 7.55 (s, 1H)	Meth 41

46		(300 MHz, CD ₃ OD) 0.75-0.85 (m, 6H), 1.05-1.3 (m, 8H), 1.4-1.6 (m, 4H), 2.2 (s, 3H), 3.25 (2H), 3.7-3.95 (m, 4H), 4.7 (ABq, 2H), 5.5 (s, 1H), 6.7 (s, 1H), 6.75-7.35 (m, 9H), 7.4 (s, 1H)	Meth 65
47		783.5	Meth 66
48		802.7	Meth 68
49		(500 MHz, CD ₃ OD) 0.82 (brt, 6H), 1.05-1.26 (m, 8H), 1.42-1.56 (m, 4H), 3.27 (brs, 2H), 3.6-3.75 (m, 2H), 4.58 (ABq, 2H), 5.41 (s, 1H), 6.73-6.82 (m, 3H), 7.0 (d, 2H), 7.05 (dd, 1H), 7.25-7.36 (m, 3H), 7.41 (brd, 2H), 7.48 (d, 1H); m/z 608.3	Ex 119

Example 50

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 ammonium salt

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 22; 150 mg, 0.30 mmol) and 2-((2'R)-2'-amino-2'-phenylethanoylamino)ethanesulphonic acid (Method 28; containing DIPEA hydrochloride, 150 mg, 0.36 mmol) was dissolved in DMF (6 ml). DIPEA (0.2 ml, 1.15 mmol) and TBTU 10 (114 mg, 0.36 mmol) were added and the mixture was stirred for 2 hours at room temperature.

The solvent was evaporated at reduced pressure. The residue was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent to give the title compound 73 mg (32%). NMR (CD_3OD) 0.75-0.85 (m, 6H), 1.0-1.3 (m, 8H), 1.3-1.6 (m, 4H), 2.16 (s, 3H), 2.85-3.0 (m, 2H), 3.24 (brs, 2H) 3.5-3.85 (m, 4H), 4.70 (ABq, 2H), 5.47 (s, 1H), 6.71 (s, 1H), 6.97 (brt, 1H), 7.11 (brd, 2H), 7.23-7.45 (m, 8H); m/z 746.2.

Example 51

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

10 ammonium salt

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 17; 49 mg, 0.10 mmol) and 2-((2'R)-2'-amino-2'-phenylethanoylamino)ethanesulphonic acid (Method 28; containing DIPEA hydrochloride; 52 mg, 0.12 mmol) was dissolved in DMF (2 ml). DIPEA (0.071 ml, 0.41 mmol) and TBTU (39 mg, 0.12 mmol) was added and the mixture was stirred for 2 hours at room temperature. The solvent was evaporated at reduced pressure. The residue was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent to give the title compound 59 mg (78%). NMR (CD_3OD) 0.74-0.90 (m, 6H), 0.98-1.3 (m, 4H), 1.35-1.67 (m, 4H), 2.16 (s, 3H), 2.85-3.02 (m, 2H), 3.23 (brs, 2H) 3.52-3.90 (m, 4H), 4.70 (ABq, 2H), 5.47 (s, 1H), 6.71 (s, 1H), 6.96 (brt, 1H), 7.09 (brd, 2H), 7.21-7.48 (m, 8H); m/z 718.4.

Example 52

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N-(carboxymethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-

25 benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N-(ethoxycarbonylmethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 72; 44 mg, 0.063 mmol) was dissolved in THF : H_2O , 1:1, (2 ml) and NaOH (1 M, 0.126 mmol) was added. The mixture was stirred at ambient temperature for 30 1 hour. The reaction mixture was acidified with HCl (1 M), diluted to 10 ml and extracted with DCM (3 x 10 ml). The combined organic layers were dried (MgSO_4) and the solvent was evaporated to give the title compound 33 mg (78%). NMR (300 MHz) 0.78-0.85 (m, 6H),

1.02- 1.70 (m, 8H), 2.20 (s, 3H), 3.15/3.21 (ABq, 2H), 3.78 (m, 2H), 3.94/4.20 (dABq, 2H), 4.64 (q, 2H), 5.91 (d, 1H), 6.65 (s, 1H), 6.98-7.52 (m, 11H), 8.17 (d, 1H).

Example 53

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N-(1"-carboxy-1"-phenylmethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

The title compound was synthesised from 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N-(1"-methoxycarbonyl-1"-phenylmethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 73) by the

10 procedure of Example 52. NMR (500 MHz) 0.76-0.84 (m, 6H), 1.05-1.73 (m, 8H), 2.14 (s, 3H), 3.16 (m, 2H), 3.74 (m, 2H), 4.48 (m, 2H), 5.53 (d, 2H), 5.96 (d, 2H), 6.63 (s, 1H), 6.97-7.48 (m, 13H), 7.86 (m, 1H), 8.17 (m, 1H).

Example 54

15 1,1-Dioxo-3-ethyl-3-butyl-5-phenyl-7-methylthio-8-[1-[N-((R)- α -carboxybenzyl)carbamoyl]ethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-{1-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoyl]ethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 10, 0.050 g, 7.6×10^{-5} mol) in DMF (4 ml) was added sodium thiomethylate (0.021 g, 3.0×10^{-4}

20 mol) and the solution was stirred for 4 hours at ambient temperature. The mixture was concentrated and the residue was partitioned between water and ether. The aqueous phase was extracted two more times with ether and the combined organic extracts were dried ($MgSO_4$), concentrated and purified by HPLC. The title compound was obtained in 0.030 g (63 %) as a white solid. NMR (CD_3OD) 0.75-0.90 (m, 6H), 1.00-1.30 (m, 4H), 1.40-1.70 (m, 7H), 2.15
25. (d, 3H), 3.10-3.30 (m, 2H), 3.55-3.95 (m, 2H), 4.80-4.95 (m, 2H), 5.30 (d, 1H), 6.70-7.50 (m, 12H); m/z 625.3.

Example 55

1,1-Dioxo-3-ethyl-3-butyl-5-phenyl-7-methylthio-8-[α -[N-((R)- α -carboxybenzyl)carbamoyl]

30 benzyloxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-ethyl-3-butyl-5-phenyl-7-methylthio-8-{ α -[N-((R)- α -carboxybenzyl)carbamoyl]benzyloxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 11; 0.018 g, 2.5×10^{-5} mol) in DMF (3 ml) was added sodium thiomethylate (0.007 g, 1.0×10^{-4}

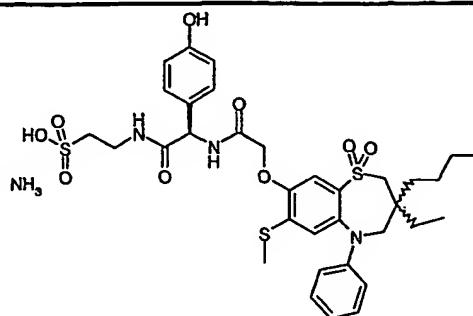
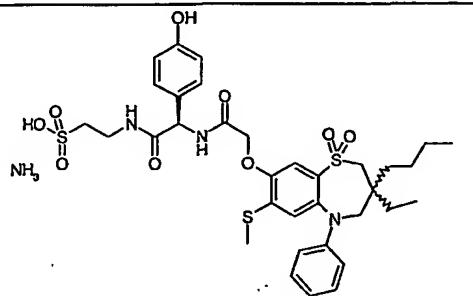
mol) and the solution was stirred for 4 hours at ambient temperature. The mixture was concentrated and the residue was partitioned between water and ether. The aqueous phase was extracted two more times with ether and the combined organic extracts were dried (MgSO_4), concentrated and purified by HPLC. The title compound was obtained in 0.015 g (89 %) as a 5 white solid. NMR (CD_3OD) 0.70-0.85 (m, 6H), 1.00-1.25 (m, 4H), 1.35-1.65 (m, 4H), 2.20 (d, 3H), 3.10-3.20 (m, 2H), 3.50-3.85 (m, 2H), 5.30 (d, 1H), 5.80 (d, 1H), 6.70 (s, 1H), 6.90-7.65 (m, 16H).

Example 56

10 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-((R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl)carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine
2-{[(2R)-2-Amino-2-(4-hydroxyphenyl)ethanoyl]amino}ethanesulphonic acid
(Method 80; 32.5 mg, 0.119 mmol) was mixed with DMF (4 ml) and N-methylmorpholine (30 μ l, 0.272 mmol). A clear solution was obtained and 1,1-dioxo-3,3-dibutyl-5-phenyl-7-15 methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 22; 50 mg, 0.099 mmol) and TBTU (38 mg, 0.119 mmol) were added successively. The reaction was stirred at ambient temperature for 30 min and the DMF was removed. The crude product was purified by preparative HPLC using an MeCN/ammonium acetate buffer (1:1). Lyophilisation gave 55 mg of the title compound (71%). NMR (500 MHz, MeOD) 0.78-0.86 (m, 6H), 1.0-20 1.3 (m, 8H), 1.4-1.6 (m, 4H), 2.15 (s, 3H), 2.85-3.00 (m, 2H), 3.25 (s, 2H), 3.55-3.68 (m, 2H), 3.75 (brs, 2H), 4.65 (ABq, 2H), 5.36 (s, 1H), 6.70 (s, 1H), 6.75 (d, 2H), 6.98 (t, 1H), 7.12 (d, 2H), 7.22 (d, 2H) 7.28 (t, 2H), 7.4 (s, 1H); m/z 762.

Examples 57-58

25 The following compounds were synthesised by the procedure of Example 56 using the appropriate starting material except that the reaction was left to proceed for 64 hours (Example 57) or 2 hours (Example 58) and the product was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (45/55 to 60/40) as eluent.

Ex	Compound	NMR (CD_3OD) and m/z	SM
57	 <p>Enantiomer 1</p>	0.75-0.84 (m, 6H) 1.00-1.27 (m, 4H), 1.37-1.64 (m, 4H) 2.14 (s, 3H), 2.86-3.00 (m, 2H), 3.22 (s, 2H), 3.53-3.68 (m, 2H), 3.85 brd, 2H), 4.68 (ABq, 2H), 5.35 (s, 1H), 6.70 (s, 1H), 6.75 (d, 2H), 6.95 (t, 1H), 7.08 (d, 2H), 7.20-7.29 (m, 4H), 7.37 (s, 1H); m/z 751 (M+NH ₄ ⁺)	Meth 23
58	 <p>Enantiomer 2</p>	0.77-0.85 (m, 6H) 1.06-1.27 (m, 4H), 1.40-1.62 (m, 4H) 2.17 (s, 3H), 2.87-3.00 (m, 2H), 3.24 (s, 2H), 3.56-3.68 (m, 2H), 3.75 (brd, 2H), 4.71 (ABq, 2H), 5.37 (s, 1H), 6.72 (s, 1H), 6.77 (d, 2H), 6.97 (t, 1H), 7.10 (d, 2H), 7.23 (d, 2H), 7.28 (t, 2H), 7.40 (s, 1H); m/z 751 (M+NH ₄ ⁺)	Meth 24

Example 59

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-carboxyethyl)

5 carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)- α -carboxybenzyl) carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38; 66.8 mg, 0.109 mmol) and β -alanine ethylester hydrochloride (23.0 mg, 0.15 mmol) was dissolved in DCM (2.5 ml) and N-methyl morpholine (0.07 ml, 0.64 mmol) was added. After stirring at ambient temperature for 5 min TBTU (45.6 mg, 0.142 mmol) was added followed by stirring for 2 hours. The reaction mixture was filtered through a short column and concentrated. The crude ester was dissolved in THF (1.5 ml) and water (1.5 ml) and NaOH (1 M, 0.20 mmol) was added. After stirring at ambient temperature for 1 hour the reaction was quenched with 1 M HCl. The reaction mixture was diluted with water (10 ml) and extracted with DCM (3 x 5 ml).

The organic layers was concentrated and purified with preparative HPLC to give the title compound (60 mg, 81%). NMR (300 MHz) 0.80 (m, 6H), 1.00-1.70 (m, 8H), 2.17 (s, 3H), 2.48 (m, 2H), 3.17 (ABq, 2H), 3.35 (m, 1H), 3.57 (m, 1H), 3.70 (m, 2H), 4.62 (ABq, 2H), 5.77 (d, 1H), 6.64 (s, 1H), 6.98 (t, 1H), 7.06 (d, 2H), 7.28 (m, 4H), 7.42 (m, 3H), 7.56 (m, 5 1H), 8.10 (m, 1H).

Examples 60-63

The following compounds were synthesised by the procedure of Example 59 using the appropriate starting material.

Ex	Compound	NMR	SM
60		0.81 (m, 6H), 1.00-1.95 (m, 10H), 2.22 (s, 3H), 3.37 (m, 2H), 3.18 (ABq, 2H), 3.48 (m, 2H), 3.75 (m, 2H), 4.66 (q, 2H), 5.75 (d, 1H), 6.67 (s, 1H), 7.00 (t, 1H), 7.09 (m, 2H), 7.20 (m, 1H), 7.30 (m, 4H), 7.44 (m, 2H), 8.25 (m, 1H)	Ex 38
61		(300 MHz, DMSO-d6) 0.74 (m, 6H), 0.95-1.50 (m, 12H), 2.16 (s, 3H), 2.28 (t, 2H), 3.24 (m, 2H), 4.74 (q, 2H), 5.33 (d, 1H), 6.69 (m, 2H), 6.85 (t, 1H), 6.99 (m, 2H), 7.16 (m, 4H), 8.33-8.45 (m, 2H)	Ex 2
62		(300 MHz) 0.81 (m, 6H), 1.00-1.74 (m, 14H), 2.22 (s, 3H), 2.31 (m, 2H), 3.10-3.35 (m, 4H), 3.73 (m, 2H), 4.62 (ABq, 2H), 5.64 (d, 1H), 6.39 (brs, 1H), 6.67 (s, 1H), 6.96-7.10 (m, 3H), 7.25-7.48 (m, 7H), 8.21 (d, 1H)	Ex 38

63		0.81 (m, 6H), 1.03-1.55 (m, 12H), 2.19 (s, 3H), 2.55 (m, 2H), 3.18 (m, 2H), 3.46 (m, 1H), 3.58 (m, 1H), 3.74 (m, 2H), 4.64 (ABq, 2H), 5.80 (m, 1H), 6.64 (s, 1H), 7.01 (t, 1H), 7.08 (d, 2H), 7.30 (m, 5H), 7.44 (m, 3H), 8.11 (m, 1H)	Ex 1
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Example 64

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-(R)-alpha-carboxy-4-methoxybenzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-{N-[{(R)-alpha-(t-butoxycarbonyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 78; 48 mg, 0.070 mmol), bromoethyl(trimethylammoniumbromide) (57 mg, 0.230 mmol), tetrabutylammonium bromide (3 mg, 0.009 mmol) and Cs₂CO₃ (71 mg, 0.22 mmol) were added to CH₃CN (1.0 ml) and the reaction mixture was heated at reflux overnight. The solvent
10 was evaporated and the residue was added to water (10 ml), extracted with DCM (3 x 5 ml) and dried (MgSO₄). The crude ester was dissolved in DCM (2.5 ml), TFA (0.3 ml) was added and the reaction mixture was stirred at room temperature overnight. The solvents were evaporated and the crude product was purified with preparative HPLC to give the title compound (23 mg, 51 %). NMR (DMSO-d₆) 0.74 (m, 6H), 0.94-1.60 (m, 8H), 2.17 (s, 3H), 3.25 (m, 2H), 3.69 (s, 3H), 4.70 (ABq, 2H), 4.95 (brs, 1H), 6.71 (s, 1H), 6.83 (m, 3H), 6.97 (d, 2H), 7.20 (m, 4H), 7.27 (s, 1H), 8.37 (brs, 1H).

15 3.25 (m, 2H), 3.69 (s, 3H), 4.70 (ABq, 2H), 4.95 (brs, 1H), 6.71 (s, 1H), 6.83 (m, 3H), 6.97 (d, 2H), 7.20 (m, 4H), 7.27 (s, 1H), 8.37 (brs, 1H).

Example 65

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-[(alpha-[N'-(2-sulphoethyl)carbamoyl]-alpha-methylbenzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine ammonium salt

20 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(alpha-carboxy-alpha-methylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 18; 27 mg, 0.041 mmol) was dissolved in DCM (2 ml). Taurine tetrabutylammonium salt (45 mg, 0.123 mmol) and TBTU (16 mg, 0.050 mmol) was added successively and the mixture was stirred for 5 hours at ambient temperature. The solvent was evaporated and the product was purified by preparative HPLC using an MeCN/ammonium acetate buffer (50/50) as eluent. Lyophilisation

25

gave the title compound in 62% yield (20 mg). NMR showed 16% of the product to remain as a tetrabutylammonium salt. NMR (500 MHz) 0.75-0.9 (m, 6H), 1.0-1.3 (m, 8H), 1.3-1.6 (m, 4H), 1.95 (s, 3H), 2.1 (s, 3H), 2.9 (brs, 2H), 3.05 (brs, 2H), 3.55 (ABd, 2H), 3.75 (brs, 2H), 4.55 (ABq, 2H), 6.6 (s, 1H), 6.9-7.6 (m, 12H), 8.2-8.3 (brs, 1H); m/z 777 (M+NH₄⁺).

5

Examples 66-67

The following compounds were synthesised by the procedure of Example 65 using the appropriate starting material.

Ex	Compound	NMR (CD ₃ OD) and M/z	SM
66		777 (M+NH ₄ ⁺)	Ex 1
67		0.75-0.85 (m, 6H), 1.02 (t, 12H), 1.05-1.3 (m, 4H), 1.3-1.7 (m, 20H), 2.17 (s, 3H), 2.85-2.99 (m, 2H), 3.19-3.26 (m, 10H), 3.52-3.92 (m, 4H), 4.71 (ABq, 2H), 5.47 (s, 1H), 6.72 (s, 1H), 6.96 (t, 1H), 7.09 (brd, 2H), 7.23-7.44 (m, 8H); m/z 735.2 (M+NH ₄ ⁺)	Ex 25

Example 68

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-[alpha-[N'-(carboxymethyl)carbamoyl]-alpha-methylbenzyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-[alpha-[N'-(methoxycarbonylmethyl)carbamoyl]-alpha-methylbenzyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

15 (Method 44; 20 mg, 0.028 mmol) was dissolved in 2.5 ml of a THF/water mixture (4/1).

LiOH (2 mg, 0.084 mmol) was added and the mixture was stirred for 1 hour at ambient temperature. The title compound was purified with preparative HPLC using an

MeCN/ammonium acetate buffer (50/50) as eluent. The MeCN was evaporated and the remaining buffer was acidified with acetic acid. Lyophilisation gave 10 mg product (51%).

NMR 0.7-0.9 (m, 6H), 1.0-1.35 (m, 8H), 1.35-1.6 (m, 4H), 2.0 (s, 3H), 2.2 (s, 3H), 3.2 (brs, 2H), 3.65-3.85 (brs, 2H), 3.9-4.1 (d, 2H), 4.5-4.7 (ABq, 2H), 6.6 (s, 1H), 6.8 (brs, 1H), 6.9-7.5 (m, 11H), 8.1 (s, 1H); m/z 727 (M + NH₄⁺).

5 **Example 69**

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-[α-[N'-(2-sulphoethyl)carbamoyl]-2-fluorobenzyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[α-carboxy-2-fluorobenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 15; 20 mg, 0.030 mmol), taurine tetrabutylammonium salt (20 mg, 0.054 mmol) and DIPEA (25 mg, 0.19 mmol) was dissolved in DMF (0.4 ml). TBTU (15 mg, 0.047 mmol) was added and the mixture was stirred for 30 min at room temperature. The product was separated from the reaction mixture by preparative HPLC using MeCN/ammonium acetate buffer (50:50) as eluent. 7 mg (29%) of the title compound was obtained. M/z = 764.5.

15

Example 70

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-(R)-{α-[N'-(R)-{α-[N''-(carboxymethyl)carbamoyl]benzyl}carbamoyl)methylcarbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

20 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(carboxymethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 43; 35 mg, 0.050 mmol) and (R)-α-[N-(t-butoxycarbonylmethyl)carbamoyl]benzylamine (Method 86; 20 mg, 0.076 mmol) were dissolved in DCM (2 ml) and 2,6-lutidine (0.03 ml, 0.26 mmol) was added. After stirring at ambient temperature for 5 min, 25 TBTU (20 mg, 0.062 mmol) was added and stirring was continued for 3 hours. The reaction mixture was filtered through a column using DCM : EtOAc; 3 : 1 as eluent. The t-butyl ester was then dissolved in DCM (6 ml) and TFA (1 ml) was added. After stirring at ambient temperature overnight the solvents were evaporated. Toluene was added and evaporated twice. No further purification was necessary to give the title compound (40 mg, 93 %). NMR 30 (500 MHz, DMSO-d₆) 0.75 (m, 6H), 0.95-1.50 (m, 12H), 2.16 (s, 3H), 3.25 (m, 2H), 3.75 (m, 2H), 3.90 (dd, 1H), 4.73/4.84 (ABq, 2H), 5.54 (m, 2H), 5.58 (d, 1H), 6.68 (s, 1H), 6.85 (t, 1H), 6.99 (d, 2H), 7.18-7.46 (m, 13H), 8.51-8.73 (m, 4H).

Example 71**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(α-carboxy-4-hydroxybenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-
5 1,5-benzothiazepine (Method 22; 61 mg, 0.12 mmol) and methyl (2S)-amino(4-hydroxyphenyl)acetate hydrochloride (31 mg, 0.14 mmol) were dissolved in DCM (4 ml) and
2,6-lutidine (0.04 ml, 0.34 mmol) was added. After stirring at room temperature for 5 min
TBTU (53 mg, 0.17 mmol) was added and stirring was continued for 2 hours. The reaction
mixture was filtered through a short column. The crude methyl ester was dissolved in THF
10 (1.5 ml) and water (1.0 ml) and NaOH (aq., 1 M, 0.39 mmol) was added. The reaction
mixture was stirred at room temperature for 8 hours, quenched with HCl (1 M) and extracted
with DCM (3 x 5 ml). The collected organic layers were concentrated and purified with
preparative HPLC using MeCN/ammonium acetate buffer (50:50) to give the title compound
(57 mg, 72 %). NMR (500 MHz, CD₃OD) 0.81 (m, 6H), 1.05-1.26 (m, 8H), 1.40-1.55 (m,
15 4H), 2.17 (s, 3H), 3.24 (brs, 2H), 3.74 (brs, 1H), 4.66 (ABq, 2H), 6.70-6.75 (m, 3H), 6.99 (t,
1H), 7.11 (d, 2H), 7.22-7.30 (m, 4H), 7.40 (s, 1H).

Example 72**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-(S)-{α-[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine ammonium salt**

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(α-carboxy-4-hydroxybenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 71; 31 mg, 0.047 mmol) and tetrabutylammonium taurine (57 mg, 0.155 mmol) were dissolved in DCM (2 ml). After stirring at room temperature for 5 min TBTU (24 mg, 0.075 mmol) was
25 added and stirring was continued for 6 hours. The solvent was evaporated and the residue was purified with preparative HPLC (twice to remove all tetrabutylammonium salt) using MeCN/ammonium acetate buffer to give the title compound 6 mg (16%). M/z 762.2.

Example 73 and Example 74

1,1-Dioxo-3-(R/S)-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-(R)-{ α -[N'-(R)-(2-imidazol-5-yl-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-(R)-1'-phenyl-1'-carboxymethyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38; 56.4 mg, 0.092 mmol) and methyl D-histidinate dihydrochloride (25.2 mg, 0.104 mmol) were added to DCM (3 ml). N-methyl morpholine (0.05 ml, 0.41 mmol) was added followed by TBTU (40 mg, 0.12 mmol). The reaction mixture was stirred at 4°C for 1 hour 30 min and at 10 room temperature for 3 hours. More TBTU (15 mg, 0.047 mmol) and DIPEA (0.025 ml, 0.14 mmol) were added and the reaction mixture was stirred at room temperature for another 30 min. The solvent was evaporated and the residue was filtered through a short column with MeOH as eluent. The crude methyl ester was dissolved in THF (1.0 ml) and water (1.0 ml) and NaOH (aq., 1 M, 0.15 mmol) was added. The reaction mixture was stirred at room 15 temperature for 2 hours and was quenched with HCl (1 M). The solvents were evaporated the residue was purified with preparative HPLC using MeCN/ammonium acetate buffer. The compound eluted as two peaks, assumed to be the two diastereomers. First peak (10 mg, 14%). Second peak (16.8 mg, 24%). First peak: NMR (DMSO-d₆) 0.74 (m, 6H), 0.95-1.60 (m, 8H), 2.17 (s, 3H), 2.82 (m, 2H), 3.23 (m, 2H), 4.27 (m, 1H), 4.80 (ABq, 2H), 5.60 (d, 1H), 6.55 (brs, 1H), 6.70 (s, 1H), 6.84 (t, 1H), 6.96 (d, 2H), 7.14-7.28 (m, 6H), 7.33 (s, 1H), 7.44 (brs, 1H), 8.54 (d, 1H), 8.60 (brs, 1H); m/z 748.4. Second peak: NMR (DMSO-d₆) 0.74 (m, 6H), 0.95-1.60 (m, 8H), 2.17 (s, 3H), 2.92 (dABq, 2H), 3.23 (m, 2H), 4.41 (m, 1H), 4.79 (ABq, 2H), 5.60 (d, 1H), 6.70 (s, 1H), 6.78 (s, 1H), 6.84 (t, 1H), 6.96 (d, 2H), 7.16-7.34 (m, 6H), 7.40 (m, 2H), 7.55 (s, 1H), 8.55 (d, 1H), 8.71 (d, 1H); m/z 748.4.

25

Example 75

1,1-Dioxo-3,3-dibutyl-5-(4-t-butylphenyl)-7-methylthio-8-(N-{(R)- α -[N'-(carboxymethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

The title compound was isolated as a byproduct in synthesis of 1,1-dioxo-3,3-dibutyl-30 5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(carboxymethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 43). Approximately 1 g of this compound was purified with preparative HPLC (MeCN/ammonium acetate buffer (50:50)) to give the title compound (32 mg). NMR (500 MHz, DMSO-d₆) 0.73 (m, 6H), 0.90-

1.40 (m, 12H), 1.24 (s, 9H), 2.16 (s, 3H), 3.23 (m, 2H), 3.65/3.75 (dABq, 2H), 4.72/4.82 (ABq, 2H), 5.60 (d, 1H), 6.65 (s, 1H), 6.97 (d, 2H), 7.23-7.35 (m, 6H), 7.45 (d, 2H), 8.58 (d, 1H), 8.62 (t, 1H).

5 **Example 76**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-(N-((R)- α -carboxybenzyl)carbamoylmethylthio)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-carboxymethylthio-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 81; 38 mg, 0.080 mmol) and D-phenylglycine methyl ester hydrochloride (24 mg, 0.12 mmol) were dissolved in DCM (2 ml) and N-methyl morpholine (0.05 ml, 0.42 mmol) was added. After stirring at room temperature for 5 min TBTU (44 mg, 0.14 mmol) was added and stirring was continued for 2 hours. The reaction mixture was filtered through a short column. The resulting product was dissolved in THF (1 ml) and water (1 ml) and NaOH (aq., 0.2 ml, 1 M) was added and the reaction mixture was stirred at room temperature for 2 hours. The reaction was quenched with HCl (1 M), diluted with water (10 ml) and extracted with DCM (3 x 3 ml). Purification with preparative HPLC yielded the title compound (40 mg, 82 %). NMR (DMSO-d₆) 0.75 (m, 6H), 0.96-1.60 (m, 8H), 3.22 (m, 2H), 3.56 (ABq, 2H), 3.89 (s, 3H), 4.81 (d, 1H), 6.78 (t, 1H), 6.83 (d, 2H), 6.89 (s, 1H), 7.11-7.23 (m, 7H), 7.31 (s, 1H), 8.37 (m, 1H).

20

Example 77

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-carboxymethylthio-8-[N-(α -carboxybenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-ethoxycarbonylmethylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 82; 21 mg, 0.038 mmol) and phenylglycine methyl ester hydrochloride (12 mg, 0.061 mmol) were dissolved in DCM (1.5 ml) and N-methyl morpholine (0.02 ml, 0.19 mmol) was added. After stirring at ambient temperature for 5 min TBTU (18 mg, 0.056 mmol) was added and stirring was continued for 2 hours. The reaction mixture was filtered through a short column. The crude diester was dissolved in THF (1 ml) and water (1 ml) and NaOH (aq., 0.1 ml, 1 M) was added. The reaction mixture was stirred at ambient temperature for 2 hours, quenched with HCl (1 M), diluted with water (10 ml) and extracted with DCM (3 x 3 ml). The collected organic layers were concentrated and purified with preparative HPLC using MeCN/ammonium acetate buffer (30:70 → 40:60) to

give the title compound (20 mg, 80%). NMR (CD_3OD) 0.80 (m, 6H), 1.03-1.26 (m, 4H), 1.38-1.65 (m, 4H), 1.96 (s, 3H), 3.20 (s, 2H), 3.44 (s, 2H), 3.67 (brs, 1H), 3.76 (brs, 1H), 4.67 (ABq, 2H), 5.29 (s, 1H), 6.89 (s, 1H), 6.92 (t, 1H), 7.05 (d, 2H), 7.19-7.32 (m, 5H), 7.41 (s, 1H), 7.45 (d, 2H).

5

Example 78

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(R)-(α-{N'-[(R)-N''-(2-hydroxy-1-carboxyethyl)carbamoylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

10 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-
(carboxymethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-
benzothiazepine (Example 43; 50 mg, 0.072 mmol), *tert*-butyl *o*-(*tert*-butyl)-D-serinate
hydrochloride (22 mg, 0.087 mmol) and N-methylmorpholine (40 mg, 0.40 mmol) were
dissolved in DCM (1 ml). TBTU (29 mg, 0.090 mmol) was added and the mixture was stirred
15 for 1 hour at room temperature. The reaction mixture was evaporated and the residue was
filtered through a short column (DCM : EtOAc, 1:4). The substance obtained (ca. 60 mg) was
dissolved in DCM (1 ml). TFA (0.59 g, 5.2 mmol) was added and the mixture was stirred for
2 hours at room temperature. The solvent was evaporated and the residue was purified by
preparative HPLC using MeCN/ammonium acetate buffer (50:50) as eluent. 38 mg (72%) of
20 the title compound was obtained. NMR (300 MHz, DMSO-d_6) 0.7-0.8 (m, 6H), 0.9-1.5 (m,
12H), 2.2 (s, 3H), 3.2-3.9 (m, 10H), 4.2 (brs, 1H), 4.8 (ABq, 2H), 5.6 (d, 1H), 6.7 (s, 1H), 6.8-
7.5 (m, 11H), 8.0 (d, 1H), 8.6 (d, 1H), 8.7 (t, 1H).

Example 79

25 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(R)-(α-{N'-[(S)-N''-(2-hydroxy-1-carboxyethyl)carbamoylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-
(carboxymethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-
30 benzothiazepine (Example 43; 50 mg, 0.072 mmol), *tert*-butyl *O*-(*tert*-butyl)-L-serinate
hydrochloride (22 mg, 0.087 mmol) and N-methylmorpholine (40 mg, 0.40 mmol) were
dissolved in DCM (1 ml). TBTU (29 mg, 0.090 mmol) was added and the mixture was stirred
for 1 h at room temperature. The reaction mixture was evaporated and the residue was filtered

through a short column (DCM : EtOAc, 1:4). The substance obtained (ca. 60 mg) was dissolved in DCM (1 ml). TFA (0.44 g, 3.9 mmol) was added and the mixture was stirred for 18 hours at room temperature. The solvent was evaporated and the residue was purified by preparative HPLC using MeCN/ammonium acetate buffer (50:50) as eluent. 33 mg (63%) of 5 the title compound was obtained. NMR (300 MHz, DMSO-d₆) 0.7-0.8 (m, 6H), 0.9-1.5 (m, 12H), 2.2 (s, 3H), 3.2-3.9 (m, 10H), 4.2 (m, 1H), 4.8 (ABq, 2H), 5.6 (d, 1H), 6.7 (s, 1H), 6.8-7.5 (m, 11H), 7.9 (d, 1H), 8.6 (d, 1H), 8.7 (t, 1H).

Example 80

10 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-[(R)-α-[N'-(1,1-dicarboxymethyl)carbamoyl]benzyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

15 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-carboxybenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38; 50 mg, 0.082 mmol), dimethylaminomalonate (60 mg, 0.13 mmol) and N-methylmorpholine (55 µl, 0.5 mmol) were dissolved in DCM (3 ml), TBTU (42 mg, 0.13 mmol) was added and the mixture was stirred for 15 mins. The solvent was evaporated under reduced pressure. The residue was dissolved in ethanol (95%) (2 ml) and a solution of NaOH (80 mg, 2 mmol) in water (80 µl) was added. The reaction mixture was stirred for 4 hours. The mixture was neutralized with 20 acetic acid. The solvent was evaporated under reduced pressure and the residue was purified by preparative HPLC using MeCN/ammonium acetate buffer (40:60) as eluent. The collected fractions were lyophilised to obtain 4 mg (7%) of the title compound. NMR (300 MHz, CD₃OD) 0.75-0.9 (m, 6H), 1.0-1.3 (m, 4H), 1.4-1.65 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.7 (brs, 2H), 4.65-4.8 (m, 2H), 5.75 (s, 1H), 6.75 (s, 1H), 6.9-7.55 (m, 11H).

25

Examples 81-87

The following compounds were synthesised by the procedure of Example 80 using the appropriate starting material except that 2,6-lutedine was used instead of N-methylmorpholine and the ration of eluent was MeCN/ammonium acetate buffer (45:55). Reaction time at each 30 stage varied slightly.

Ex	Compound	NMR (300 MHz, CD ₃ OD)	SM
81		(500 MHz) 0.8-0.95 (m, 6H), 1.05-1.35 (m, 4H), 1.4-1.7 (m, 4H), 2.2 (s, 3H), 3.25 (s, 2H), 3.7-3.9 (m, 4H), 4.4-4.45 (m, 1H), 4.7-4.8 (m, 2H), 5.7 (s, 1H), 6.75 (s, 1H), 6.95-7.6 (m, 11H)	Ex 38
82		0.75-0.9 (m, 6H), 1.05-1.3 (m, 4H), 1.4-1.6 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.7-3.95 (m, 4H), 4.25-4.3 (m, 1H), 4.75 (ABq, 2H), 5.65 (s, 1H), 6.75 (s, 1H), 7.95-7.55 (m, 11H)	Ex 38
83		0.75-0.9 (m, 6H), 1.05-1.35 (m, 8H), 1.4-1.6 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.7-3.9 (m, 4H), 4.35-4.45 (m, 1H), 4.7 (ABq, 2H), 5.7 (s, 1H), 6.75 (s, 1H), 6.95-7.55 (m, 11H)	Ex 1
84 1		0.75-0.9 (m, 6H), 1.05-1.3 (m, 8H), 1.4-1.6 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.7-3.9 (m, 4H), 3.3-3.4 (m, 1H), 4.7 (ABq, 2H), 5.65 (s, 1H), 6.7 (s, 1H), 6.95-7.55 (m, 11H)	Ex 1
85 2		0.75-0.9 (m, 6H), 1.05-1.3 (m, 8H), 1.4-1.6 (m, 4H), 3.25 (s, 3H), 3.6-3.9 (m, 6H), 4.3-4.5 (m, 2H), 4.7 (ABq, 2H), 5.65 (s, 1H), 6.7 (s, 1H), 6.95-7.5 (m, 11H)	Ex 83

86 3		0.75-0.9 (m, 6H), 1.05-1.3 (m, 8H), 1.4-1.6 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.6-3.9 (m, 6H), 4.35-4.5 (m, 2H), 4.7 (ABq, 2H), 5.6 (s, 1H), 6.7 (s, 1H), 6.95-7.55 (m, 11H)	Ex 83
87 4		0.75-0.9 (m, 6H), 1.05-1.3 (m, 8H), 1.4-1.6 (m, 4H), 2.2 (d, 3H), 3.15-3.35 (m, 5H), 3.5-3.85 (4H), 4.4-4.5 (m, 1H), 4.6-4.7 (m, 2H), 5.6 (s, 1H), 6.7 (s, 1H), 6.95-7.55 (m, 11H)	Ex 1

¹ Eluent ratio (55:45); ² Eluent ratio - variable gradient; ³ Eluent ratio (50:50); ⁴ Eluent ratio (60:40)

Example 88

5 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(alpha-carboxybenzyl) carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(alpha-methoxycarbonylbenzyl) carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 87; 55.2 mg, 0.064 mmol) was dissolved in THF (2 ml) and 0.5 ml water. LiOH (3.1 mg, 0.127 mmol) was added 10 and the mixture was stirred for 1 hour. Water (1 ml) was added and the mixture was acidified with 0.1M HCl and extracted with DCM (3 x 2 ml). The DCM phase was dried with and concentrated. The solid product was co-evaporated with diethyl ether and dissolved in HPLC grade water. Lyophilisation gave the title compound as a white solid in 68% yield (28 mg).

NMR 0.77-0.85 (m, 6H), 1.03-1.25 (m, 8H), 1.34-1.57 (m, 4H), 2.16 (s, 3H), 3.18 (brs, 2H), 15 3.75 (brs, 2H), 4.65 (ABq, 2H), 5.7 (d, 1H), 6.63 (s, 1H), 7.0 (t, 1H), 7.1 (d, 2H), 7.26-7.48 (m, 8H), 7.85 (d, 1H); m/z 639.

Example 89

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-{(S)- α -[N'-(carboxymethyl)carbamoyl]benzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-{(S)- α -[N'-
5 (methoxycarbonylmethyl)carbamoyl]benzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-
benzothiazepine (Method 88; 19 mg, 0.027 mmol) was hydrolysed by LiOH (1.3 mg, 0.054
mmol) in THF (1 ml) and water (0.3 ml). After 1 hour water (3 ml) was added and the mixture
acidified using 0.1M HCl and extracted with DCM (3 x 3 ml). The organic layer was dried
and evaporated yielding 16 mg (82% yield) of the title compound. NMR 0.77-0.85 (m, 6H),
10 1.0-1.3 (m, 8H), 1.34-1.57 (m, 4H), 2.17 (s, 3H), 3.18 (s, 2H), 3.75 (brs, 2H), 3.90-4.20 (m,
2H), 4.65 (ABq, 2H), 5.87 (m, 1H), 6.63 (s, 1H), 6.98-7.50 (m, 12H), 8.12-8.20 (m, 1H); m/z
696.

Example 90

15 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-{(S)- α -[N'-(2-sulphoethyl)carbamoyl]benzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine sodium salt

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(α-carboxybenzyl)
carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 88; 41 mg, 0.064
mmol) was dissolved in 3 ml DCM. Taurine tetrabutylammonium salt (70 mg, 0.191 mmol)
20 and TBTU (25 mg, 0.078 mmol) were added successively and the mixture was stirred
overnight at ambient temperature. The solvent was evaporated and the product was purified
by preparative HPLC using an MeCN/ammonium acetate buffer gradient (45/55 to 55/45) as
eluent. Lyophilisation of the collected fractions and then ion-exchange chromatography over
4 g Amberlite CG 120, Na⁺-form, gave the title compound in 85% yield (42 mg). NMR 0.7-
25 0.8 (m, 6H), 0.9-1.2 (m, 8H), 1.3-1.5 (m, 4H), 2.0 (s, 3H), 2.9-3.2 (m, 2H+2H), 3.3-3.8 (m,
2H+2H), 4.4-4.7 (m, 2H), 5.6 (m, 1H), 6.57 (s, 1H), 6.9-7.5 (m, 11H), 7.8-8.1 (m, 2H); m/z
746.

Example 91

30 The following compound was synthesised by the procedure of Example 90 using the
appropriate starting material except the product was purified using a buffer gradient of 40/60
to 70/30 and then lyophilised to give the ammonium salt.

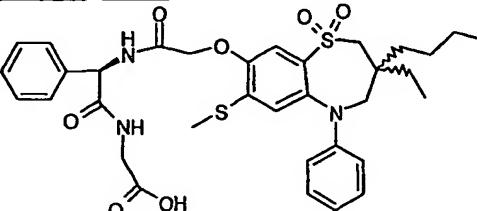
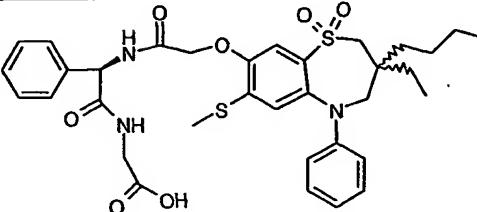
Ex	Compound	NMR (CD_3OD) and m/z	SM
91		0.76-0.84 (m, 6H), 1.03-1.27 (m, 8H), 1.38-1.55 (m, 4H), 2.15 (s, 3H), 2.95 (t, 2H), 3.24 (s, 2H), 3.58 (dt, 2H), 3.75 (brs, 2H), 3.85 (ABdd, 2H), 4.72 (ABq, 2H), 5.51 (s, 1H), 6.70 (s, 1H), 6.97 (t, 1H), 7.11 (d, 2H), 7.25-7.40 (m, 6H), 7.46 (d, 2H); m/z 803	Ex 43

Example 92**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-[N-(R)-α-[N'-(carboxymethyl)carbamoyl]****5 benzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine sodium salt**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-carbamoylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine sodium salt (WO 01/66533; 120 mg, 0.278 mmol) dissolved in DCM (4 ml) was added to a solution of α -[N-(t-butoxycarbonylmethyl)carbamoyl]benzylamine (Method 86; 90%, 150 mg, 0.511 mmol) in DCM (3 ml). 2,6-Dimethylpyridine (65 \square l, 0.559 mmol) and TBTU (137 mg, 0.427 mmol) were added and the reaction mixture was stirred at ambient temperature overnight. The solution was filtered over using DCM/EtOAc (8/2) as eluent. The solvent was evaporated. DCM (4 ml) and TFA (0.6 ml) were added and the mixture was stirred overnight. The solvent was evaporated and the crude product was purified by preparative HPLC on a Chromasil C₁₈ column. An MeCN/ammonium acetate buffer gradient (50/50 to 100/0) was used as mobile phase. The MeCN was evaporated and lyophilisation gave the title compound in 36% yield (62 mg). NMR 0.73-0.82 (m, 6H), 1.00-1.23 (m, 4H), 1.30-1.65 (m, 4H), 3.05-3.18 (m, 2H), 3.65 (brs, 2H), 3.75 (ABdd, 2H), 4.46 (ABq, 2H), 5.70 (d, 1H), 6.79-7.24 (m, 10H), 7.36 (d, 2H), 7.46 (d, 1H), 7.83 (d, 1H), 8.00 (brs, 1H); m/z 622.

20 Example 93-94

The following compounds were synthesised by the procedure of Example 92 using the appropriate starting material except that the HPLC-chromatography was performed on a Chromasil C₈ column and the eluent gradient was 45/55 to 60/40.

Ex	Compound	NMR (CD_3OD) and m/z	SM
93	 <p>Enantiomer 1</p>	0.75-0.84 (m, 6H) 1.00-1.27 (m, 4H), 1.38-1.66 (m, 4H) 2.15 (s, 3H), 3.22 (s, 2H), 3.75 (brs, 2H), 3.83 (ABdd, 2H), 4.69 (ABq, 2H), 5.60 (s, 1H), 6.71 (s, 1H), 6.96 (t, 1H), 7.09 (d, 2H), 7.23-7.37 (m, 5H), 7.39 (s, 1H), 7.46 (d, 2H); m/z 668	Meth 23; Meth 86
94	 <p>Enantiomer 2</p>	0.78-0.85 (m, 6H) 1.04-1.27 (m, 4H), 1.41-1.65 (m, 4H) 2.17 (s, 3H), 3.24 (s, 2H), 3.68 (brs, 2H), 3.89 (ABdd, 2H), 4.72 (ABq, 2H), 5.62 (s, 1H), 6.73 (s, 1H), 6.97 (t, 1H), 7.11 (d, 2H), 7.26-7.38 (m, 5H), 7.41 (s, 1H), 7.48 (d, 2H); m/z 668	Meth 24; Meth 86

Example 95

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-[(R)- α -(N'-[2-[(ethoxy)methyl]phosphoryl]ethyl)carbamoyl]benzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 2-[(methyl)(ethyl)phosphoryl]ethylamine (Helv.Chim.Acta; GE; 75; 8; 1992; 2545-2552; 16 mg, 0.106 mmol) in DCM (2 ml) was added at 0 °C 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38; 50 mg, 0.082 mmol) DIPEA (42 mg, 0.328 mmol) and TBTU (34 mg, 0.106 mmol) under argon. The reaction mixture was stirred at room temperature for 110 min and then DCM was added and the solution washed with NaHCO_3 (aq, sat) and brine. The organic layer was dried and the solvent evaporated under reduced pressure. The residue purified by chromatography and the product eluted with DCM/methanol (100:5). Yield 43 mg (71%). NMR (500 MHz) 0.78-0.85 (m, 6 H), 1.02-1.54 (m, 12H), 1.6-1.75 (br, 1H), 1.8-2.10 (m, 3H), 2.21 (s, 3H), 3.10-3.25 (m,

2H), 3.51-3.84 (m, 4 H), 3.9-3.99 (m, 1H), 4.01-4.09 (m, 1H), 4.54-4.69 (dd, 2H), 5.51 (d, 1H), 6.68 (s, 1 H), 6.96-7.02 (m, 1H), 7.03-7.18 (m, 3H), 7.25-7.42 (m, 6H), 7.43-7.48 (m, 2H), 8.05-8.15 (m, 1H).

5 Examples 96-97

The following compounds were synthesised by the procedure of Example 95 using the appropriate starting material.

Ex	Compound	NMR and m/z	SM
96		0.76-0.85 (m, 6H), 1.00-1.52 (m, 12H), 1.55-1.75 (m, 1H), 1.95-2.12 (br, 1H), 2.20 (s, 3H), 3.10-3.25 (m, 2H), 3.55-3.85 (m, 4H), 3.85-4.00 (m, 2H), 4.03-4.13 (m, 2H), 4.6 (q, 2H), 5.64 (d, 1H), 6.66 (s, 1H), 7.78 (br, 1H), 6.95-7.10 (m, 3H), 7.23-7.40 (m, 6H), 7.43-7.49 (m, 2H), 8.07 (d, 1H); m/z 760.3	Ex. 38
97 1		(600 MHz) 0.75-0.82 (m, 6H), 1.0-1.42 (m, 13H), 1.64 (brs, 1H), 2.18 (s, 3H), 3.08-3.24 (m, 2H), 3.50-3.84 (m, 4H), 3.87-4.13 (m, 2H), 4.54-4.68 (m, 2H), 5.56-5.62 (m, 1H), 6.63 (s, 1H), 6.87-7.10 (m, 3H), 7.24-7.40 (m, 7H), 7.43-7.49 (m, 2H), 7.98-8.05 (m, 1H); m/z 730.5	Ex 38

¹ Amine: Tetrahedron; EN; 49; 47; 1993; 11055-11064

10 Example 98

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-{N-[R]-alpha-(N'-(2-[(hydroxy)(methyl)phosphoryl]ethyl)carbamoyl]benzyl}carbamoylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-{N-[R]-alpha-(N'-(2-[(ethoxy)(methyl)phosphoryl]ethyl)carbamoyl]benzyl}carbamoylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 95; 27 mg, 0.036 mmol) in ethanol (1.5 ml) was

added at 0°C 2 M aqueous NaOH (0.22 ml, 0.44 mmol). The reaction mixture was stirred at room temperature for 24 hours. Acetic acid (0.2 ml) was added. The solvent was evaporated under reduced pressure and the residue was extracted with DCM/water. The DCM layer was separated, washed with brine, dried and evaporated under reduced pressure. Recrystallization

5 of the residue from DCM/ether/petroleum ether gave the title compound 23 mg (89%). NMR (600 MHz) 0.74-0.82 (m, 6H), 1.0-1.70 (m, 11H), 1.90-2.09 (m, 2H), 2.16 (s, 3H), 3.05-3.24 (m, 2H), 3.40-3.85 (m, 4H), 4.50-4.65 (dd, 2H), 5.55 (d, 1H), 6.63 (s, 1H), 6.93-7.07 (m, 3H), 7.20-7.50 (m, 9H), 8.10 (d, 1H); m/z 716.3.

10 **Example 99**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[(hydroxy)(ethoxy)phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[(diethoxy)phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 96; 13 mg, 0.017 mmol) in MeCN (0.5 ml) was added 1 M aqueous LiOH (0.171 ml, 0.171 mmol) dropwise. The reaction mixture was stirred at room temperature for 3 days. Acetic acid was added and the solvent evaporated under reduced pressure. The crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (45:55) as eluent to give the title compound 11 mg (88%). NMR (600 MHz, CD₃OD) 0.77-0.84 (m, 6H), 1.00-1.30 (m, 7H), 1.40-1.65 (m, 4H), 2.17 (s, 3H), 3.23 (brs, 2H), 3.51 (d, 2H), 3.6-3.85 (m, 4H), 4.70 (dd, 2H), 5.57 (s, 1H), 6.72 (s, 1H), 6.96 (t, 1H), 7.09 (d, 2H), 7.25-7.31 (m, 3H), 7.32-7.36 (m, 2H), 7.40 (s, 1H), 7.45 (d, 2H); m/z 732.4.

25 **Example 100**

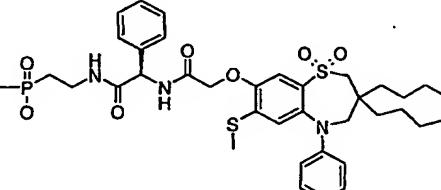
1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[(hydroxy)(methyl)phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[(ethoxy)(methyl)phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 97; 85 mg, 0.12 mmol) in MeCN (2.4 ml) was added at 0°C 1 M aqueous LiOH (1.17 ml, 1.17 mmol) dropwise. The reaction mixture was stirred at room temperature for 20 hours. Acetic acid was added and the solvent evaporated

under reduced pressure. The crude product was purified by column chromatography using DCM/MeOH/Et₃N (100:15:0.2 and 100:30:0.2) as eluent to give the title compound 62 mg (76%). NMR (CD₃OD) 0.75-0.84 (m, 6H), 1.0-1.70 (m, 11H), 2.15 (s, 3H), 3.22 (brs, 2H), 3.35 (d, 2H), 3.60-3.90 (m, 2H), 4.70 (dd, 2H), 3.55 (s, 1H), 6.71 (s, 1H), 6.96 (t, 1 H), 7.09 5 (d, 2H), 7.23-7.38 (m, 5H), 7.40 (s, 1H), 7.46 (d, 2H); m/z 702.3

Example 101

The following compound was synthesised by the procedure of Example 100 using the appropriate starting material.

Ex	Compound	NMR (600 MHz, CD ₃ OD) and m/z	SM
101		0.76-0.83 (m, 6H), 1.05-1.55 (m, 15H), 1.91-1.99 (m, 2H), 2.15 (s, 3H), 3.24 (brs, 2H), 3.40-3.50 (m, 2H), 3.66-3.86 (m, 2H), 4.69 (dd, 2H), 5.42 (s, 1H), 6.70 (s, 1H), 6.92 (t, 1H), 7.11 (d, 2H), 7.25-7.39 (m, 6H), 7.43 (d, 2H); m/z 744.3	Ex 104

10

Example 102

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[di-(t-butoxy)phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

15 To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl) carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38; 80 mg, 0.131 mmol) and di-(t-butoxy)phosphorylmethyamine (Tet. Lett.; EN; 33; 1; 1992; 77-80; 37 mg, 0.164 mmol) in DCM (5 ml) was added 2,6-lutidine (28 mg, 0.262 mmol) and TBTU (53 mg, 0.164 mmol). The reaction mixture was stirred at room 20 temperature for 2 hours and 50 min. The solvent was evaporated under reduced pressure and the crude product was purified by column chromatography using DCM/MeOH (100:4) as eluent to give the title compound 92 mg (86%). NMR (500 MHz) 0.77-0.86 (m, 6H), 1.03-1.75 (m, 26H), 2.22 (s, 3H), 2.10-2.25 (m, 2H), 3.45-3.90 (m, 4H), 4.61 (dd, 2H), 5.52 (d, 1H), 5.94 (brs, 1H), 6.67 (s, 1H), 7.0 (t, 1H), 7.07 (d, 2H), 7.26-7.48 (m, 8H), 8.12 (d, 1H); 25 m/z 704.22 [M-2(t-butyl)+2H].

Example 103

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[di-(hydroxy) phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-α-{N'-[di-(*t*-butoxy) phosphorylmethyl]carbamoyl}benzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 102; 72 mg, 0.088 mmol) in DCM (4 ml) was added at 0°C TFA (1 ml). The reaction mixture was stirred at room temperature for 2 hours. The solvent was evaporated under reduced pressure and the residue was extracted with DCM/water. The
10 organic layer was separated, washed with brine, dried and evaporated under reduced pressure. The residue was suspended in ether and the crystals filtered to give the title compound 60 mg (97%). NMR (500 MHz, DMSO-d₆) 0.70-0.80 (m, 6H), 0.99-1.61 (m, 8H), 2.18 (s, 3H), 2.80-4.0 (m, 6H), 4.80 (dd, 2H), 5.65 (d, 1H), 6.71 (s, 1H), 6.80-7.02 (m, 3H), 7.15-7.35 (m, 6H), 7.48 (d, 2H), 8.50-9.20 (m, 2H); m/z 704.3

15

Example 104

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-[(R)-α-(N'-{2-[(methyl)(ethyl) phosphoryl]ethyl}carbamoyl)benzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

20 To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 1; 60 mg, 0.094 mmol) and 2-[(methyl)(ethyl)phosphoryl]ethylamine (Helv.Chim.Acta; GE; 75; 8; 1992; 2545-2552; 20 mg, 0.132 mmol) was added at 0°C 2,6-lutidine (20 mg, 0.19 mmol) and TBTU (39 mg, 0.121 mmol) under argon. The reaction mixture was stirred at room
25 temperature for 70 min and then DCM was added and the solution washed with water and brine. The organic layer was dried and the solvent evaporated under reduced pressure. The residue was purified by column chromatography using DCM/MeOH (100:5) as eluent to give the title compound 67 mg (92%). NMR (300 MHz) 0.74-86 (m, 6H), 1.0-1.60 (m, 18H), 1.80-2.05 (m, 2H), 2.20 (s, 3H), 2.17 (s, 2H), 3.47-3.80 (m, 4H), 3.88-4.10 (dd, 2H), 5.52 (d, 1H),
30 6.65 (s, 1H), 6.95-7.12 (m, 3H), 7.13-7.42 (m, 7H), 7.43-7.49 (m, 2H), 8.05-8.16 (m, 1H); m/z 772.4.

Example 105

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-[(R)- α -[N'-(2-mercaptopro-1-carboxyethyl)carbamoyl]benzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-[(R)- α -[N'-(2-mercaptopro-1-carboxyethyl)carbamoyl]benzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 91; 37 mg, 0.036 mmol) in DCM (1 ml) was added at 0°C TFA (1 ml) under argon blanketing. The ice-bath was removed and triethylsilane (42 mg, 0.36 mmol) was added. The reaction mixture was stirred at room temperature for 2 hours and then the solvent evaporated under reduced pressure. The crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (40:60 to 60:40) as eluent to give the title compound 16 mg (59%). NMR (500 MHz, CD₃OD) 0.76-0.85 (m, 6H), 1.05-1.60 (m, 12H), 2.17 (s, 3H), 2.77-2.92 (m, 2H), 3.24 (brs, 2H), 3.61-3.88 (m, 2H), 4.56 (t, 1H), 4.70 (dd, 2H), 5.65 (s, 1H), 6.71 (s, 1H), 6.98 (t, 1H), 7.12 (d, 2H), 7.25-7.43 (m, 6H), 7.50 (d, 2H); m/z 742.4.

15

Example 106

The following compound was synthesised by the procedure of Example 105 using the appropriate starting material.

Ex	Compound	NMR (500 MHz, CD ₃ OD) and m/z	SM
106		0.77-0.85 (m, 6H), 1.03-1.28 (m, 8H), 1.38-1.58 (m, 4H), 2.15 (s, 3H), 2.87-3.5 (m, 2H), 3.25 (s, 2H), 3.75 (brs, 2H), 4.55 (s, 1H), 4.71 (dd, 2H), 5.66 (s, 1H), 6.71 (s, 1H), 6.98 (t, 1H), 7.12 (d, 2H), 7.25-7.43 (m, 6H), 7.50 (d, 2H); m/z 742.28	Meth 93

20 **Example 107**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-(2-[N-[(R)- α -(carboxy)benzyl]carbamoyl]ethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-(2-[N-[(R)- α -(carboxy)benzyl]carbamoyl]ethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 90; 77 mg, 0.108 mmol) in DCM (3 ml) was added at 0°C TFA (0.75 ml). The reaction

mixture was stirred at room temperature for 2 h and 45 min. The solvent was evaporated under reduced pressure and the crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (40:60 to 50:50) as eluent to give the title compound 60 mg (82%). NMR (500MHz, CD₃OD) 0.75-0.85 (m, 6H), 1.0-1.25 (m, 4H), 1.40-1.64 (m, 4H), 5 2.75-2.90 (m, 2H), 3.26 (s, 2H), 3.50-3.90 (m, 2H), 4.30-4.41 (m, 2H), 5.43 (s, 1H), 6.99 (t, 1H), 7.05-7.13 (m, 3H), 7.23-7.34 (m, 5H), 7.45 (d, 2H), 7.52 (s, 1H); m/z 658.

Example 108

The following compound was synthesised by the procedure of Example 107 using the 10 appropriate starting material.

Ex	Compound	NMR (500 MHz, CD ₃ OD) and m/z	SM
108		0.78-0.85 (m, 6H), 1.02-1.30 (m, 8H), 1.38-1.58 (m, 4H), 1.87 (s, 3H), 2.15 (s, 3H), 2.77-2.83 (m, 1H), 2.87-2.94 (m, 1H), 3.24 (s, 2H), 3.74 (brs, 2H), 4.53-4.59 (m, 1H), 4.68 (dd, 2H), 5.66 (s, 1H), 6.71 (s, 1H), 6.98 (t, 1H), 7.12 (d, 2H), 7.25-7.31 (m, 3H), 7.32-7.36 (m, 2H), 7.40 (s, 1H), 7.49 (d, 2H); m/z 756.23	Meth 92

Example 109

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[{(R)-alpha-(N'-(2-[(methyl)(ethyl)phosphoryl]ethyl)carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine

15 benzothiazepine

To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-((R)- α -carboxy-4-hydroxybenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 2; 80 mg, 0.122 mmol) and 2-[(methyl)(ethyl)phosphoryl]ethylamine (Helv.Chim.Acta; GE; 75; 8; 1992; 2545-2552; 24 mg, 0.159 mmol) in DCM (2 ml) was added 2,6-lutidine (26 mg, 0.244 mmol) and TBTU (51 mg, 0.159 mmol) under argon. The reaction mixture was stirred at room temperature for 60 min, then diluted with DCM. The solution was washed with water, brine, dried and the solvent was evaporated under reduced pressure. The residue was purified by column chromatography using DCM/MeOH (100:7) as eluent to give the title compound

67 mg (92%). NMR (600 MHz), 0.74-0.80 (m, 6H), 1.0-1.55 (m, 18H), 1.82-1.98 (m, 2H), 2.15 (s, 3H), 3.14 (brs, 2H), 3.40-3.56 (m, 2H), 3.70 (brs, 2H), 3.89-4.02 (m, 2H), 4.51 (dd, 2H), 5.33 (t, 1H), 6.61 (s, 1H), 6.65-6.72 (m, 2H), 6.95 (t, 1H), 7.03 (d, 2H), 7.12-7.19 (m, 3H), 7.22-7.26 (m, 2H), 7.32 (s, 1H), 8.11 (t, 1H); m/z 788.56.

5

Example 110

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[*(R*)- α -(*N'*-{2-[(methyl)(hydroxy)phosphoryl]ethyl}carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine

10 To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[*(R*)- α -(*N'*-{2-[(methyl)(ethyl)phosphoryl]ethyl}carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 104; 37 mg, 0.047 mmol) in MeCN/MeOH (4 ml, 1:1) was added 1 M aqueous LiOH (0.8 ml, 0.8 mmol). The reaction mixture was stirred at room temperature for 40 min. Acetic acid was added and the solvent evaporated under reduced pressure. The crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (40:60 and 45:55) as eluent to give the title compound 35 mg (96%). NMR (500 MHz, CD₃OD) 0.78-0.85 (m, 6H), 1.06-1.28 (m, 11H), 1.39-1.57 (m, 4H), 1.72-1.85 (m, 2H), 2.16 (s, 3H), 2.24 (s, 2H), 3.40-3.50 (m, 2H), 3.65-3.84 (m, 2H), 4.69 (dd, 2H), 5.36 (s, 1H), 6.71 (s, 1H), 6.76 (d, 2H), 6.99 (t, 1H), 7.13 (d, 2H), 7.22-7.33 (m, 4H), 20 7.39 (s, 1H); m/z 760.27

Example 111

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-[*(R*)- α -[*(R*)-*N'*-(2-methylsulphinyl-1-carboxyethyl)carbamoyl]benzyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

25 The title compound was separated as by-product from the synthesis of Example 108. NMR (500 MHz, CD₃OD) 0.78-0.85 (m, 6H), 1.02-1.60 (m, 12H), 2.16 (d, 3H), 2.53 (d, 3H), 3.08-3.18 (m, 1H), 3.24 (s, 2H), 3.35 (v br, 1H), 3.75 (v br, 2H), 4.62 (v br, 1H), 4.71 (dd, 2H), 5.60 (d, 1H), 7.71 (s, 1H), 6.98 (t, 1H), 7.12 (d, 2H), 7.25-7.42 (m, 6H), 7.47 (d, 2H); m/z 772.25.

30

Example 112

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[(S)-N'-(3-methylthio-2-carboxypropyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[(S)-N'-(3-methylthio-2-methoxycarbonylpropyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 94; 68 mg, 0.087 mmol) in ethanol (5 ml) was added NaOH (9 mg in 0.4 ml water) at 0°C. The reaction mixture was stirred at room temperature for 2.5 hours. Acetic acid was added and the solvent evaporated under reduced pressure. The
10 crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (40:60 to 60:40) as eluent to give the title compound 52 mg (76%). NMR (500 MHz, CD₃OD) 0.79-0.86 (m, 6H), 1.05-1.29 (m, 8H), 1.40-1.58 (m, 4H), 1.84-1.93 (m, 4H), 2.01-2.21 (m, 5H), 2.26-2.34 (m, 1H), 3.26 (s, 2H), 3.76 (brs, 2H), 4.52-4.58 (m, 1H), 4.70 (dd, 2H), 5.61 (s, 1H), 6.73 (s, 1H), 7.0 (t, 1H), 7.14 (d, 2H), 7.27-7.43 (m, 6H), 7.49 (d, 2H); m/z 770.16.

15

Example 113

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[(S)-N'-(2-methylthio-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[(S)-N'-(2-mercaptop-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 106; 15 mg, 0.02 mmol) in methanol (1.5 ml) was added sodium methoxide (0.104 mmol in 0.14 ml methanol) and methyl iodide (0.16 mmol) under nitrogen. The reaction mixture was stirred at room temperature for 50 min. Acetic acid was added. The solvent was evaporated under reduced pressure and the residue was extracted with
20 DCM/water. The organic layer was separated, washed with brine, dried and evaporated under reduced pressure to give the title compound 4 mg (26%). NMR (500 MHz, CD₃OD) 0.75-8.30 (m, 6H), 1.03-1.57 (m, 12H), 2.10 (s, 3H), 2.17 (s, 3H), 2.83-2.30 (m, 1H), 3.0-3.25 (m, 1H), 3.26 (s, 2H), 3.77 (brs, 2H), 4.58-4.63 (m, 1H), 4.72 (dd, 2H), 5.64 (s, 1H), 6.72 (s, 1H), 7.0 (t, 1H), 7.12 (d, 2H), 7.28-7.52 (m, 8H); m/z 756.25.

30

Example 114**1,1-Dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-[N-{(R)- α -[N'-(carboxymethyl)carbamoyl]benzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-[N-{(R)- α -[N'-(t-butoxycarbonylmethyl)carbamoyl]benzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 102; 129 mg, 0.164 mmol) in DCM (5 ml) was added at 0°C TFA (1.5 ml) under nitrogen. The reaction mixture was stirred at room temperature for 2 hours. The solvent was evaporated under reduced pressure and the crude product was purified by preparative HPLC using MeCN and ammonium acetate buffer (40:60 to 50:50) as eluent to give the title compound 77 mg (63%). NMR (500 MHz, CD₃OD) 0.84 (t, 6H), 1.10-1.22 (m, 8H), 1.35-1.45 (m, 4H), 2.34 (s, 3H), 3.19-3.27 (m, 2H), 3.55 (s, 2H), 3.87 (dd, 2H), 4.67 (dd, 2H), 5.61 (s, 1H), 7.09-7.15 (m, 3H), 7.27-7.37 (m, 6H), 7.47 (d, 2H); m/z 748.03 (M+NH₃).

Example 115**1,1-Dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-[N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 118; 0.050 g, 0.105 mmol) in DMF (4 ml) was added 2-{[(2R)-2-amino-2-(4-hydroxyphenyl)ethanoyl]amino}ethanesulphonic acid (Method 80; 0.037 g, 0.135 mmol) and N-methylmorpholine (0.040 ml, 0.363 mmol). The mixture was stirred for 10 min and then was TBTU (0.044 g, 0.137 mmol) added. The reaction mixture was stirred for two days before the solvent was removed under reduced pressure. The residue was purified by preparative HPLC using a MeCN/ammonium acetate buffer to give the title compound in 0.042 g (55 %) as a white solid. NMR (DMSO-d₆) 0.60-0.80 (m, 6H), 1.05-1.50 (m, 8H), 2.15 (s, 3H), 2.45-2.55 (m, 2H), 3.05-3.80 (m, 6H), 4.70 (ABd, 1H), 4.80 (ABd, 1H), 5.25 (d, 1H), 6.65-6.75 (m, 3H), 6.80-7.05 (m, 3H), 7.10-7.25 (m, 4H), 7.30 (s, 1H), 8.20-8.30 (m, 1H), 8.45 (d, 1H).

Example 116**1,1-Dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-[N-{(R)- α -[N'-(carboxymethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-carboxymethoxy-
5 2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 118; 0.050 g, 0.105 mmol) in DCM (4 ml)
was added (R)- α -[N-(*t*-butoxycarbonylmethyl)carbamoyl]benzylamine (Method 86; 0.036 g,
0.136 mmol) and *N*-methylmorpholine (0.040 ml, 0.363 mmol). The mixture was stirred for 5
min and then was TBTU (0.044 g, 0.137 mmol) added. The reaction mixture was stirred for
two days and then was TFA (1.5 ml) added. After 1.5 h, the solution was diluted with toluene,
10 before the solvent was removed under reduced pressure. The residue was purified by
preparative HPLC using a MeCN/ammonium acetate buffer to give the title compound in
0.020 g (29 %) as a white solid. NMR (DMSO-d₆) 0.60-0.80 (m, 6H), 1.05-1.50 (m, 8H), 2.15
(s, 3H), 3.10-3.80 (m, 6H), 4.70 (ABd, 1H), 4.85 (ABd, 1H), 5.60 (d, 1H), 6.70 (s, 1H), 6.80-
7.05 (m, 3H), 7.15-7.50 (m, 8H), 8.35 (brs, 1H), 8.55 (d, 1H).

15

Example 117**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-[N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-carboxymethoxy-
20 2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 6; 0.020 g, 4.09*10⁻⁵ mol) in DMF (4 ml)
was added 2-{[(2*R*)-2-amino-2-(4-hydroxyphenyl)ethanoyl]amino}ethanesulphonic acid
(Method 80; 0.014 g, 5.10*10⁻⁵ mol) and *N*-methylmorpholine (0.020 ml, 1.81*10⁻⁴ mol). The
mixture was stirred for 10 min and then was TBTU (0.016 g, 4.98*10⁻⁵ mol) added. The
reaction mixture was stirred for 3 h and then the solvent was removed under reduced pressure.
25 The residue was purified by preparative HPLC using a MeCN/ammonium acetate buffer to
give the title compound in 0.023 g (75 %) as a white solid. NMR (500 MHz, DMSO-d₆) 0.65-
0.80 (m, 6H), 0.80-1.50 (m, 12H), 2.40-2.60 (m, 2H), 3.15-3.45 (m, 4H), 3.60 (s, 3H), 3.65
(brs, 2H), 4.60 (ABd, 1H), 4.70 (ABd, 1H), 5.25 (d, 1H), 6.50 (s, 1H), 6.70-7.25 (m, 10H),
7.35 (s, 1H), 8.20-8.30 (m, 1H), 8.50 (d, 1H), 9.40 (brs, 1H).

30

Example 118**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-8-[N-[(R)- α -[N'-(2-sulphoethyl)carbamoyl]4-hydroxybenzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 115; 0.020 g, 4.63×10^{-5} mol) in DMF (4 ml) was added 2-{[(2R)-2-amino-2-(4-hydroxyphenyl)ethanoyl]amino}ethanesulphonic acid (Method 80; 0.017 g, 6.20×10^{-5} mol) and N-methylmorpholine (0.016 ml, 1.46×10^{-4} mol). The mixture was stirred for 10 min and then TBTU (0.019 g, 5.92×10^{-5} mol) was added. The reaction mixture was stirred overnight and then the solvent was removed under reduced pressure. The residue was purified by preparative HPLC using a MeCN/ammonium acetate buffer to give the title compound in 0.008 g (24 %) as a white solid. NMR (500 MHz, DMSO-d₆) 0.65-0.80 (m, 6H), 0.80-1.60 (m, 8H), 2.40-2.55 (m, 2H), 3.20-3.40 (m, 4H), 3.65 (brs, 2H), 4.65 (ABd, 1H), 4.70 (ABd, 1H), 5.25 (d, 1H), 6.65-7.45 (m, 13H), 8.20-8.30 (m, 1H), 8.60 (d, 1H), 9.40 (brs, 1H).

15

Example 119**1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-[N-(α -(R)-carboxybenzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

The title compound was synthesized from 1,1-dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-[N-(α -(R)-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 45) by the procedure of Method 109. NMR (CD₃OD) 0.81 (brt, 6H), 1.03-1.3 (m, 8H), 1.32-1.59 (m, 13H), 3.24 (brs, 2H), 3.57-3.77 (m, 2H), 4.61 (brs, 2H), 5.51 (s, 1H), 6.83 (d, 1H), 7.0-7.1 (m, 3H), 7.26-7.43 (m, 7H), 7.49 (d, 1H); m/z 708.5.

25

Example 120**1,1-Dioxo-3,3-dibutyl-5-[4-(N'-t-butylureido)phenyl]-8-[N-(α -(R)-carboxybenzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3,3-dibutyl-5-(4-(N'-t-butylureido)phenyl)-8-[N-(α -(R)-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 111; 30 mg, 0.042 mmol) was dissolved in THF (1.5 ml), H₂O (0.5 ml) and LiOH (42 mg, 0.064 mmol, monohydrate) was added. The mixture was stirred for 2 hours. The compound was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95

to 100/0) as eluent to give the title product, 24 mg (82%). NMR (CD_3OD) 0.81 (brt, 6H), 1.05-1.26 (m, 8H), 1.35 (s, 9H), 1.38-1.57 (m, 4H), 3.25 (brs, 2H), 3.6-3.77 (m, 2H), 4.61 (ABq, 2H), 5.45 (s, 1H), 6.84 (d, 1H), 7.01-7.11 (m, 3H), 7.24 (d, 2H), 7.26-7.37 (m, 3H), 7.37-7.42 (m, 2H), 7.50 (d, 1H); m/z 707.5.

5

Preparation of Starting Materials

The starting materials for the Examples above are either commercially available or are readily prepared by standard methods from known materials. For example, the following reactions are an illustration, but not a limitation, of some of the starting materials used in the 10 above reactions.

Method 1

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[1'-(ethoxycarbonyl)ethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

15 Sodium carbonate (0.30 g, 2.83 mmol), 2-bromopropanoic acid ethyl ester (0.145 g, 0.796 mmol) and tetrabutylammonium bromide (0.022 g, 0.07 mmol) was added to a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 96/16051; 0.300 g, 0.663 mmol) in MeCN (10 ml). The suspension was heated under reflux overnight. The solvent was evaporated and the crude mixture was 20 extracted (DCM/H₂O), dried (MgSO₄), evaporated and purified by flash chromatography (Hex:EtOAc - 5:1) to give the title compound 0.346 g (95 %) as a white solid. NMR 0.70-0.85 (m, 6H), 1.00-1.75 (m, 8H), 1.35 (t, 3H), 1.70 (d, 3H), 3.05-3.25 (m, 2H), 3.55-3.90 (m, 2H), 4.20-4.35 (m, 2H), 4.80 (q, 1H), 7.00-7.10 (m, 3H), 7.15 (s, 1H), 7.25-7.35 (m, 2H), 7.45 (s, 1H).

25

Method 2

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[1'-carboxyethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

30 Sodium hydroxide (0.045 g, 1.13 mmol) was added to a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[1-(ethoxycarbonyl)ethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 1; 0.050 g, 0.090 mmol) in EtOH (4 ml, 95 %) and heated under reflux. After 1.5 hours AcOH (0.2 ml) was added and most of the solvent was removed under reduced pressure. The crude product was extracted (DCM/H₂O), dried (MgSO₄) and

evaporated to give the title compound 0.031 g (65 %) as white solid. NMR (500 MHz, CD₃OD) 0.70-0.85 (m, 6H), 0.95-1.25 (m, 4H), 1.35-1.70 (m, 4H), 2.65 (d, 3H), 3.10-3.35 (m, 2H), 3.45-3.95 (m, 2H), 4.70 (q, 1H), 6.90-7.35 (m, 6H), 7.45 (s, 1H).

5 Method 3

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[1'-phenyl-1'-ethoxycarbonylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

Ethyl α-bromophenylacetate (0.139 g), Na₂CO₃ (0.200 g) and tetrabutylammonium bromide (0.034 g) were added to a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 96/16051; 0.200 g, 0.442 mmol) in MeCN (6 ml). The suspension was heated under reflux overnight before the solvent was removed under reduced pressure. The crude product was extracted (DCM/H₂O) and purified by flash chromatography (Hex:EtOAc-5:1) to give the title compound 0.256 g (94 %) as a white solid. NMR 0.65-0.85 (m, 6H), 0.95-1.65 (m, 8H), 3.00-3.15 (m, 2H), 3.50-3.80 (m, 2H), 3.70-3.80 (2s, 3H), 5.60 (s, 1H), 5.65 (d, 1H) 7.00-7.60 (m, 17H), 8.05-8.20 (2d, 1H).

Method 4

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[1'-phenyl-1'-carboxymethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

Lithium hydroxide (0.019 g) was added to a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[1'-phenyl-1'-ethoxycarbonylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 3; 0.244 g, 0.397 mmol) in THF/H₂O (2/1, 3 ml). After 2 days the solvent was removed under reduced pressure and the crude mixture was purified by HPLC to give the title compound 0.215 g (92 %) as a white solid. NMR (CD₃OD) 0.60-0.80 (m, 6H), 0.90-1.25 (m, 4H), 1.30-1.60 (m, 4H), 3.05-3.30 (m, 2H), 3.40-3.90 (m, 2H), 5.55 (s, 1H), 6.85-7.70 (m, 12H).

Method 5

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-

30 benzothiazepine

Ethyl bromoacetate (0.13 ml), Na₂CO₃ (0.40 g) and tetrabutylammonium bromide (0.030 g) were added to a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (synthesised by the of WO9616051 for the

corresponding 3-butyl-3-ethyl analogue; 0.400 g, 0.927 mmol) in MeCN (10 ml). The suspension was heated under reflux overnight before most of the solvent was removed under reduced pressure. The crude product was extracted (DCM/H₂O) and filtered through a short silica-column (DCM:EtOAc-1:4) to give the title compound 0.476 g (99 %). NMR 0.65-0.85
5 (m, 6H), 0.95-1.65 (m, 8H), 3.00-3.15 (m, 2H), 3.50-3.80 (m, 2H), 3.70-3.80 (s, 3H), 5.60 (s, 1H), 5.65 (d, 1H) 7.00-7.60 (m, 17H), 8.05-8.20 (d, 1H).

Method 6

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-
10 benzothiazepine

Lithium hydroxide (0.062 g) was added to a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 5; 0.448 g, 0.865 mmol) in THF/H₂O (2/1, 6 ml). After 1 hour AcOH (0.5 ml) was added and most of the solvent was removed under reduced pressure. The crude product was
15 purified by HPLC (MeCN) to give the title compound 0.408 g (96 %) as a white solid. NMR (CD₃OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.55 (m, 4H), 3.20 (s, 2H), 3.65 (s, 3H), 3.70 (brs, 2H), 4.50 (s, 2H), 6.50 (s, 1H), 6.90-7.30 (m, 5H), 7.40 (s, 1H).

Method 7

20 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methoxy-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-
1,5-benzothiazepine

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methoxy-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 9616051; 1.0 g), ethyl bromoacetate (0.50 g), sodium carbonate (1.2 g) and tetrabutylammonium bromide (60 mg) in MeCN (15 ml) were refluxed overnight. The
25 solvent was removed under reduced pressure and the residue was extracted (DCM/H₂O). The organic layer was separated and the solvent was removed under reduced pressure. The residue purified by chromatography (DCM/EtOAc (90:10)) to give the title compound 1.2 g (98%).
NMR (CD₃OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.55 (m, 4H), 3.20 (s, 2H), 3.65 (s, 3H), 3.70 (brs, 2H), 4.50 (s, 2H), 6.50 (s, 1H), 6.90-7.30 (m, 5H), 7.40 (s, 1H).

Method 8**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

A mixture of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 96/16051; 0.3 g), ethyl bromoacetate (0.14 g), sodium carbonate (0.3 g), tetrabutylammonium bromide (0.02 g) in MeCN (10 ml) were refluxed for 4 hours. The solvent was removed under reduced pressure. The residue was partitioned between DCM/H₂O and the organic layer was separated. The solvent was evaporated and the residue was purified by chromatography (DCM/EtOAc, 90:10) to give the title compound 0.34 g (95%). NMR (500 MHz) 0.7-0.9 (m, 6H), 1.0-1.8 (m, 11H), 3.2 (m, 2H), 3.6-3.8 (brs, 2H), 4.3 (q, 2H), 4.7 (s, 2H), 7.0-7.1 (m, 3H), 7.15 (s, 1H), 7.3 (m, 2H), 7.4 (s, 1H).

Method 9**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 8; 0.34 g) and sodium hydroxide (0.3 g) were dissolved in ethanol and the mixture was heated to reflux for 1 hour. Acetic acid (1 ml) was added and the solvent was removed at reduced pressure. The residue was partitioned between DCM/H₂O and the organic layer was separated and dried. Trituration of the residue with *n*-hexane gave the title compound 0.29 g (90%) as a solid. NMR (500 MHz) 0.7-0.8 (m, 6H), 1.0-1.7 (m, 8H), 3.1-3.2 (m, 2H), 3.6 (brs, 2H), 4.6 (s, 2H), 6.9-7.1 (m, 4H), 7.2 (m, 2H), 7.5 (s, 1H).

25 Method 10**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methoxy-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methoxy-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 7; 1.2 g) was dissolved in ethanol (20 ml). Sodium hydroxide (0.5) dissolved in H₂O (1 ml) was added and the reaction mixture was warmed to 40°C for 30 min. Acetic acid (1 ml) was added and the solvent was removed at reduced pressure. The residue was partitioned between DCM/H₂O and the organic layer was separated and dried. Trituration of the residue with *n*-hexane gave the title compound 1.1 g (97%) as a

solid. NMR 0.75-0.85 (m, 3H), 0.9 (t, 3H), 1.0-1.7 (m, 8H), 3.2 (q, 2H), 3.65 (s, 3H), 3.65-3.85 (m, 2H), 4.7 (s, 2H), 6.4 (s, 1H), 7.0 (t, 1H), 7.1 (d, 2H), 7.3 (t, 2H), 7.5 (s, 1H).

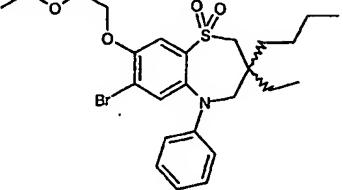
Method 11

5 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-bromo-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (synthesised by the of WO9616051 for the corresponding 3-butyl-3-ethyl analogue; 2.0 g, 4.16 mmol), ethyl bromoacetate (0.84 g, 5.03 mmol), sodium carbonate (2.0 g, 18.9 mmol) and tetrabutylammonium bromide (80 mg, 0.25 mmol) were added to MeCN (20 ml). The mixture was refluxed for 2 hours and then evaporated under reduced pressure. The residue was extracted with DCM/water. The DCM layer was separated and evaporated under reduced pressure. The residue was purified by column chromatography . The product was eluted with DCM / EtOAc (90:10) to give the title compound 2.2 g (93%). NMR 0.7-0.8 (m, 6H), 1.0-1.6 (m, 15H), 3.2 (brs, 2H), 3.7 (brs, 2H), 4.3 (q, 2H), 4.7 (s, 2H), 7.0-7.3 (m, 6H), 7.4 (s, 1H).

Methods 12-13

The following compounds were synthesised by the procedure of Method 11 using the appropriate acid and amine (source not indicated where commercially available).

Meth	Compound	M/z	SM
12	 Enantiomer 1	538	Meth 83

13	 Enantiomer 2	538	Meth 84
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Method 14**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

5 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-bromo-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 11; 2.2 g, 3.88 mmol) was dissolved in ethanol (15 ml). NaOH (0.8 g in 1.5 ml water) was added to the solution and the mixture was stirred for 30 min at room temperature. Acetic acid (2 ml) was added. The solvent was evaporated under reduced pressure and the residue was extracted with EtOAc/water. The EtOAc layer was separated, dried and evaporated under reduced pressure to give the title compound 2.0 g (95%). NMR (500 MHz) 0.7-0.8 (m, 6H), 1.0-1.5 (m, 12H), 3.2 (brs, 2H), 3.7 (brs, 2H), 4.7 (s, 2H), 7.0-7.3 (m, 6H), 7.4 (s, 1H).

Method 15**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-isopropoxy-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

15 To isopropyl alcohol (12 ml) was added sodium (115 mg, 5 mmol) and the temperature was then raised to 80°C to let the alcohol salt form. After all the sodium was dissolved 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 9; 100 mg, 0.2 mmol) was added in one portion. The reaction was then refluxed overnight, and then cooled to room temperature and quenched with acetic acid. The solvent was then removed under reduced pressure and the residue was dissolved in water and MeCN (70/30) and partially purified by HPLC. The residue was dissolved in ethylene glycol and NaOH (500 mg) was added. This reaction mixture was heated to 125°C overnight and then cooled to room temperature and quenched with acetic acid, and EtOAc (100 ml) was added. The ethylene glycol was removed by washing the organic layer with

acidic water three times. The organic layer was then concentrated and the residue was purified again as above to give the title compound 40 mg (41 %). NMR (300 MHz) 0.7-1.0 (m, 6H), 1.0-1.8 (m, 15H), 3.2 (q, 2H), 3.75 (m, 2H), 4.3 (m, 1H), 4.6 (s, 2H), 6.35 (s, 1H), 6.95-7.2 (m, 3H), 7.2-7.4 (m, 2H), 7.55 (s, 1H).

5

Method 16

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-10 1,5-benzothiazepine (Method 25; 500 mg, 1.2 mmol) was added MeCN (30 ml), tetrabutylammonium bromide (30 mg, 0.08 mmol), anhydrous sodium carbonate (500 mg, 4.7 mmol), ethyl bromoacetate (0.14 ml, 1.26 mmol) and caesium carbonate (20 mg, 0.06 mmol). This reaction mixture was then stirred overnight at 80°C. Then the solvent was removed under reduced pressure, water and DCM were added and the aqueous phase was extracted 15 three times with DCM. The combined organic phases were then dried, concentrated and purified by flash chromatography [DCM : EtOAc, 1:0, 9:1] to give the title compound 600 mg (99%). NMR (300 MHz) 0.8-1.0 (m, 6H), 1.0-1.8 (m, 11H), 2.2 (s, 3H), 3.2 (q, 2H) 3.75 (brq, 2H), 4.3 (q, 2H), 4.75 (s, 1H), 6.7 (s, 1H), 6.95 (t, 1H), 7.05 (d, 2H), 7.25 (t, 2H), 7.3 (s, 1H).

20

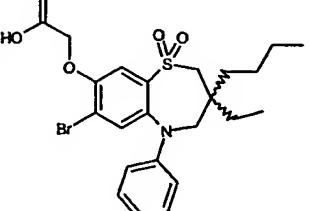
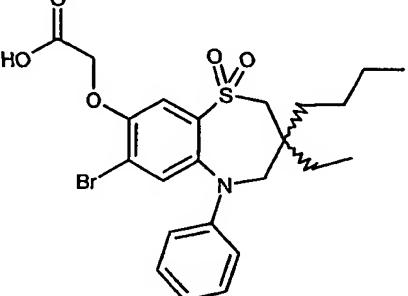
Method 17

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 16; 478 mg, 0.95 mmol) was added THF (15 ml), water (3 ml) and LiOH (34 mg, 1.4 mmol). The reaction was then stirred for 1 hour. Then acetic acid (0.2 ml) was added along with water (10 ml) and DCM (10 ml). The aqueous layer were then extracted three times with DCM. The combined organic phases were then dried and concentrated to give the title compound 450 mg (99%). NMR 0.7-0.9 (m, 6H), 1.0-1.7 (m, 8H), 2.2 (s, 3H), 3.2 (q, 2H), 3.7 (m, 2H), 4.8 (s, 2H), 6.65 (s, 1H), 6.95 (t, 1H), 7.05 (d, 2H), 30 7.25 (t, 2H), 7.35 (s, 1H), 8.4 (brs, 1H).

Method 18-19

The following compounds were synthesised by the procedure of Method 17 using the appropriate acid and amine (source not indicated where commercially available) except two equivalents of LiOH was used and the extraction was performed after 2 hours reaction time
 5 using EtOAc.

Meth	Compound	M/z	SM
18	 Enantiomer 1	510	Meth 12
19	 Enantiomer 2	510	Meth 13

Method 20

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-mesyl-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

10 To 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-ethoxycarbonylmethoxy-
 2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 16; 122 mg, 0.24 mmol) was added DCM (3 ml), water (3 ml) and potassium carbonate (120 mg, 0.87 mmol). The reaction mixture was then cooled to 0°C and m-chloroperoxybenzoic acid (160 mg, 0.51 mmol) was added in one portion. After 5 hours the reaction was quenched with DCM and saturated sodium hydrogen
 15 carbonate solution the aqueous phase were then extracted three times with DCM. The combined organic phases were dried, concentrated and purified by flash chromatography [DCM : EtOAc, 9:1] to give the title compound 46 mg (35%). NMR 0.7-0.8 (m, 6H), 1.0-1.65

(m, 11H), 3.2 (q, 2H), 3.3 (s, 3H), 3.7 (brs, 1H), 4.25 (q, 2H), 4.8 (s, 2H), 7.0-7.1 (m, 3H), 7.2-7.3 (m, 2H), 7.5 (s, 2H).

Method 21

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-mesyl-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-mesyl-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 20; 46 mg, 0.085 mmol) was added THF (5 ml), water (1 ml) and LiOH (10 mg, 0.4 mmol). The reaction was stirred for 1 hour and then 10 excess acetic acid was added to quench the reaction. Water and DCM were added and the aqueous phase was extracted three times with DCM. The combined organic phases were dried and concentrated to give the title compound 40 mg (91%). NMR 0.7-0.85 (m, 6H), 1.0-1.7 (m, 8H), 3.2 (m, 2H), 3.3 (s, 3H), 3.8 (s, 2H), 4.9 (s, 2H), 5.0 (brs, 1H), 7.05-7.15 (m, 3H), 7.3-7.4 (t, 2H), 7.5 (s, 1H), 7.6 (s, 1H).

15

Method 22 (Preparation 1)

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 14; 500 mg, 0.93 mmol) was dissolved in DMF (10 ml). Sodium methanethiolate (200 mg, 2.85 mmol) was added and the mixture was stirred for 2 hours at 50°C. Acetic acid (0.4 ml) was added and the solvent was evaporated under reduced pressure. The residue was extracted with EtOAc/water. The EtOAc layer was separated, dried and evaporated under reduced pressure to give the title compound 450 mg (96%). NMR (300 MHz) 0.7-0.8 (m, 6H), 1.0-1.6 (m, 12H), 2.2 (s, 2H), 3.2 (brs, 2H), 3.7 (brs, 2H), 4.8 (s, 2H), 6.6 (s, 1H), 6.9-7.1 (m, 3H), 7.2-7.4 (m, 3H).

Method 22 (Preparation 2)

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-

30 benzothiazepine

A solution of NaOH (4.67 g, 116 mmol) in water (10 ml) was added to a solution of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-ethoxycarbonyl-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 114; 15.45 g, 28.71 mmol) in EtOH (160 ml). The solution was

stirred for 30 min at room temperature. The solvent was removed under reduced pressure and the residue was partitioned between EtOAc and 1.0 M HCl. The aqueous layer was extracted twice more with EtOAc and the combined organic extracts were washed with brine and concentrated to give the title compound (14.28 g, 98 %) as a white powder. NMR (500 MHz,

5 DMSO-d₆) 0.65-0.80 (m, 6H), 0.90-1.50 (m, 12H), 2.20 (s, 3H), 3.25 (s, 2H), 3.65 (bs, 2H),
4.80 (s, 2H), 6.70 (s, 1H), 6.80-7.30 (m, 6H), 13.20 (s, 1H).

Method 23-24

The following compounds were synthesised by the procedure of Method 22

10 (Preparation 1) using the appropriate acid and amine (source not indicated where commercially available) except that the reactions were performed at ambient temperature and in Method 24 for extended reaction time.

Meth	Compound	M/z	SM
23	 Enantiomer 1	478	Meth 18
24	 Enantiomer 2	478	Meth 19

Method 25

15 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO9616051; 600 mg, 1.29 mmol) were added DMF (5 ml) and sodium

methanethiolate (450 mg, 6.42 mmol). The reaction was then heated to 60°C for 1 hour. The oil bath was then heated to 120°C for 4 hours. To quench the reaction, the temperature was lowered to room temperature and excess acetic acid was added quickly. The reaction was kept under a flow of nitrogen through sodium hypochlorite for 2 hours. Water and EtOAc were 5 added and the aqueous phase was extracted three times with EtOAc. The combined organic phases were washed with water, dried and concentrated under reduced pressure. The residue was then purified by flash chromatography [DCM : EtOAc, 9:1] to give the title compound 0.5 g (92%). NMR 0.65-0.8 (m, 6H), 0.95-1.6 (m, 8H), 3.1 (q, 2H), 3.6 (brq, 2H), 6.75 (s, 1H), 6.8 (t, 1H), 6.9 (d, 2H), 7.15 (t, 2H), 7.55 (s, 1H).

10

Method 26**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To 1,1-dioxo-3,3-dibutyl-5-phenyl-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-15 benzothiazepine (synthesised by the of WO9616051 for the corresponding 3-butyl-3-ethyl analogue; 40 mg, 0.08 mmol) was added DMF (2 ml), sodium methanethiolate (60 mg, 0.85 mmol) and sodium borohydride (60 mg, 1.6 mmol). The reaction was run overnight at 60°C. Additional sodium borohydride (60 mg, 1.6 mmol) and sodium methanethiolate(60 mg, 0.85 mmol) was added and the temperature was raised to 120°C. The reaction heated at this 20 temperature for 4 hours and then cooled to room temperature. Then acetic acid was added under a flow of nitrogen, through sodium hypochlorite overnight. Water and EtOAc was added and the aqueous phase was extracted three times with EtOAc. The combined organic phases were washed with HCl (1M), dried and concentrated under reduced pressure. The residue was then purified by flash chromatography [EtOAc : heptane, 1:4] to give the title 25 compound 0.34 g (93%). NMR 0.7-0.9 (m, 6H), 1.0-1.6 (m, 12H), 2.2 (s, 3H), 3.1 (s, 2H), 3.4 (s, 2H), 3.7 (brs, 2H), 6.7 (s, 1H), 6.85-7.05 (m, 2H), 7.2-7.4 (m, 2H).

Method 27**2-[*(2'R)-2'-(*t*-Butoxycarbonylamino)-2'-phenylethanoylamino]ethanesulphonic acid***30 **ammonium salt**

2-Aminoethanesulphonic acid (740 mg, 5.91 mmol) and (2R)-2-(*t*-butoxycarbonylamino)-2-phenylacetic acid (1.09 g, 4.34 mmol) were dissolved in DMF (20 ml). DIPEA (2.8 ml, 16.1 mmol) and TBTU (1.53 g, 4.78 mmol) were added and the mixture

was stirred for 2 hours at 60°C. The solvent was evaporated at reduced pressure. The residue was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent to give the title compound 589 mg (32%). NMR (CD₃OD) 1.43 (s, 9H), 2.85-3.0 (m, 2H), 3.53-3.68 (m, 2H), 5.1 (brs, 1H), 7.25-7.45 (m, 5H).

5

Method 28**2-((2'R)-2'-amino-2'-phenylethanoylamino)ethanesulphonic acid ammonium salt**

2-[(2'R)-2'-(*t*-Butoxycarbonylamino)-2'-phenylethanoylamino]ethanesulphonic acid ammonium salt (Method 27; 589 mg, 1.57 mmol) was dissolved in EtOAc (20 ml) and the mixture was cooled in an ice bath. Hydrogen chloride gas was bubbled through the reaction, the ice bath was removed and the reaction was allowed to stand for 30 minutes at room temperature. The solvent was evaporated at reduced pressure. The residue was then redissolved in EtOAc (20 ml) and cooled in an ice bath. Hydrogen chloride gas was again bubbled through the reaction, the ice bath was removed and the reaction was allowed to stand for 30 minutes at room temperature. The solvent was evaporated at reduced pressure. DIPEA in DCM was added and the mixture was evaporated at reduced pressure. This was repeated twice. The mixture was lyophilised to give the title compound 563 mg (85%) containing 1 equivalent of di-isopropylethylammoniumchloride. NMR (D₂O) 1.35-1.38 (m, 15H), 2.96-3.12 (m, 2H), 3.21 (q, 2H), 3.50-3.80 (m, 4H), 5.11 (brs, 1H), 7.45-7.55 (m, 5H).

20

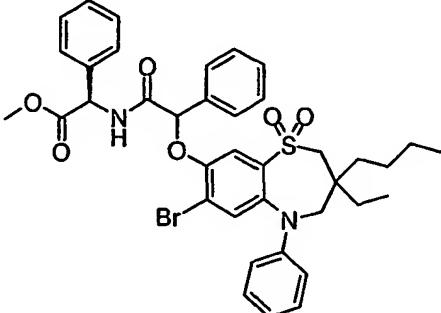
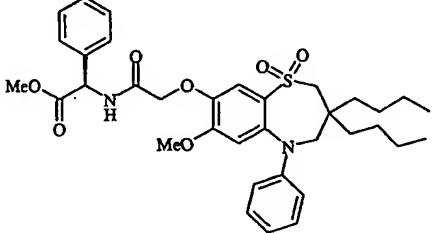
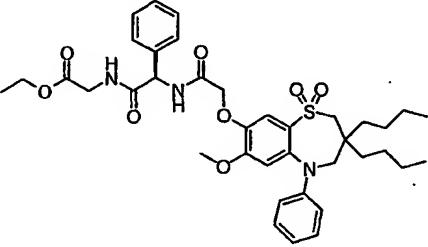
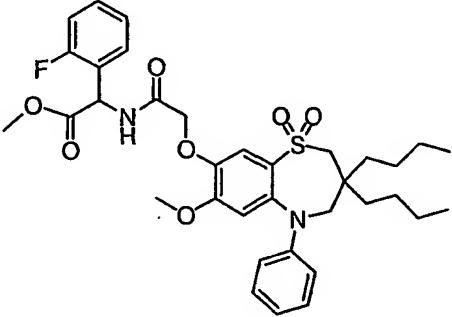
Method 29**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(R)-1'-phenyl-1'-methoxycarbonylmethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 22; 250 mg, 0.49 mmol), (R)-2-phenylglycine methyl ester hydrochloride (120 mg, 0.60 mmol) and DIPEA (300 mg, 2.3 mmol) were dissolved in DCM (10 ml) and the mixture was stirred for 5 min in room temperature. TBTU (210 mg, 0.65 mmol) was added and the mixture was stirred for 30 min at room temperature. The solvent was evaporated under reduced pressure and the residue was placed on a column and the product was eluted with DCM/EtOAc (90:10) to give the title compound 306 mg (95%). NMR (500 MHz) 0.7-0.8 (m, 6H), 1.0-1.6 (m, 12H), 2.1 (s, 3H) 3.2 brs, 2H), 3.6-3.8 (m, 5H), 4.6 (ABq, 2H), 5.6 (d, 1H), 6.6 (s, 1H), 6.9-7.5 (m, 11H), 7.9 (d, 1H).

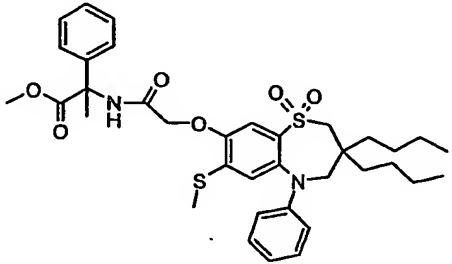
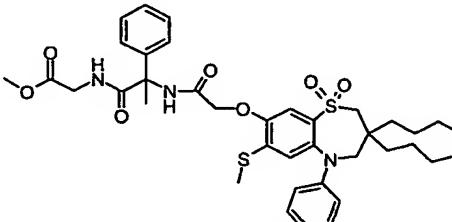
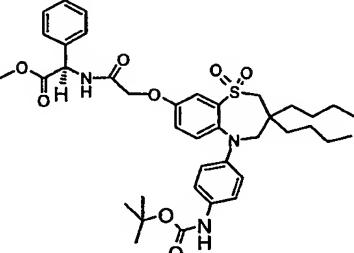
Methods 30-45

The following compounds were synthesised by the procedure of Method 29 using the appropriate acid and amine (source not indicated where commercially available) except that the reaction time was extended to 2 hours for some methods.

Meth	Compound	NMR or m/z	SM
30		(300 MHz, CD ₃ OD) 0.8-0.9 (m, 6H), 1.1-1.6 (m, 12H), 2.2 (s, 3H), 3.3 (s, 2H), 3.75 (brs, 5H), 4.7-4.8 (m, 2H), 5.45 (s, 1H), 6.7 (s, 1H), 6.8-7.3 (m, 9H), 7.45 (s, 1H)	Meth 22
31		(300 MHz, CD ₃ OD) 0.75-0.95 (m, 6H), 1.0-1.6 (m, 12H), 2.1 (s, 3H), 3.2 (s, 2H), 3.7 (s, 5H), 4.65 (s, 2H), 5.85 (s, 1H), 6.7 (s, 1H), 6.9-7.4 (m, 9H)	Meth 22
32		(300 MHz, CD ₃ OD) 0.75-0.9 (m, 6H), 1.0-1.6 (m, 12H), 2.2 (s, 3H), 3.2 (s, 2H), 3.75 (s, 5H), 4.7 (brs, 2H), 5.7 (s, 1H), 6.7 (s, 1H), 6.9-7.4 (m, 6H), 7.55-7.8 (m, 4H)	Meth 22 and Meth 71
33		0.70-0.85 (m, 6H), 0.95-1.75 (m, 8H), 1.55-1.75 (2d, 3H), 3.05-3.30 (m, 2H), 3.55-3.90 (m, 2H), 3.70-3.80 (2s, 3H), 4.75-4.90 (2q, 1H), 5.60 (d, 1H), 7.00-7.55 (m, 12H), 7.80-7.95 (m, 1H)	Meth 2

34		0.65-0.85 (m, 6H), 0.95-1.65 (m, 8H), 3.00-3.15 (m, 2H), 3.50-3.80 (m, 2H), 3.70-3.80 (2s, 3H), 5.60 (s, 1H), 5.65 (d, 1H) 7.00-7.60 (m, 17H), 8.05-8.20 (2d, 1H)	Meth 4
35		(CD ₃ OD) 0.75-0.85 (m, 6H), 1.00-1.30 (m, 8H), 1.35-1.55 (m, 4H), 3.20 (s, 2H), 3.55 (s, 3H), 3.70 (s, 3H), 3.75 (brs, 2H), 4.60 (ABq, 2H), 5.55 (s, 1H), 6.50 (s, 1H), 6.95-7.40 (m, 10H), 7.50 (s, 1H)	Meth 6
36		707.4	Ex 12
37		0.75-0.85 (m, 6H), 1.00-1.60 (m, 12H), 3.20 (s, 2H), 3.60 (s, 3H), 3.75 (brs, 2H), 3.75 (s, 3H), 4.55 (ABq, 2H), 5.85 (d, 1H), 6.40 (s, 1H), 6.95-7.45 (m, 9H), 7.55 (s, 1H), 8.05 (d, 1H)	Meth 6

38		0.75-0.85 (m, 6H), 1.00-1.60 (m, 12H), 2.20 (s, 3H), 3.20 (s, 2H), 3.75 (brs, 2H), 3.80 (s, 3H), 4.60 (ABq, 2H), 5.90 (d, 1H), 6.65 (s, 1H), 6.95-7.45 (m, 10H), 7.95 (d, 1H)	Meth 22
39		(500 MHz) 0.7-0.8 (m, 6H), 1.0-1.5 (m, 12H), 3.2 (m, 2H), 3.7-3.8 (m, 5H), 4.6 (ABq, 2H), 5.6 (d, 1H), 6.8-7.4 (m, 10H), 7.5 (s, 1H)	Meth 14
40		(300 MHz) 0.7-0.8 (m, 6H), 1.0-1.6 (m, 12H), 3.2 (brs, 2H), 3.7-3.8 (m, 5H), 4.6 (ABq, 2H), 5.6 (d, 1H), 6.8 (d, 2H), 7.0-7.4 (m, 8H), 7.9 (d, 1H)	Meth 14
41		766.4 (M-(t-butyl)+2H)	Ex 12
42		739.3	Ex 38

43		667	Meth 22
44		724	Ex 18
45		722.5	Meth 109

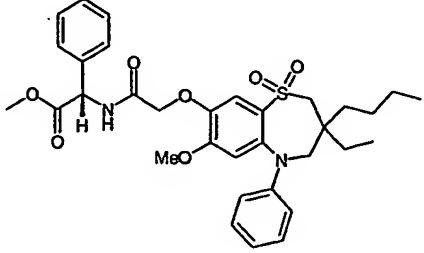
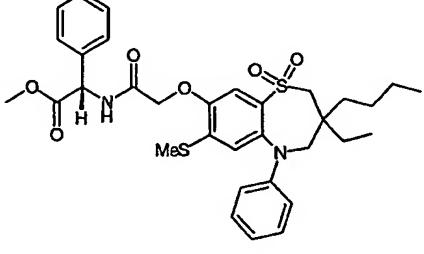
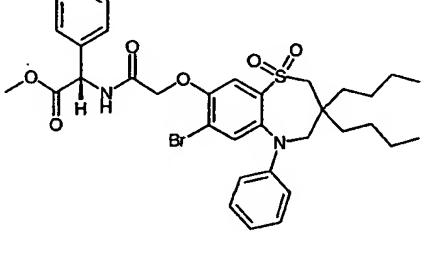
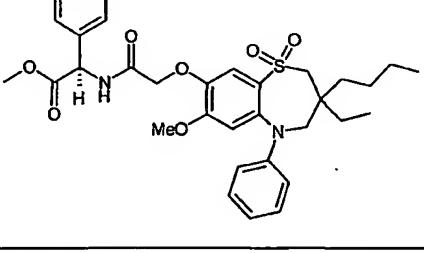
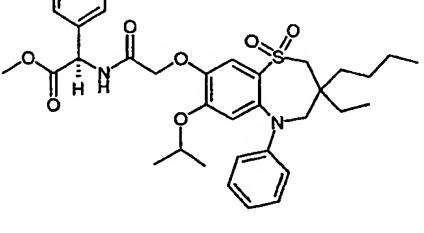
Method 46

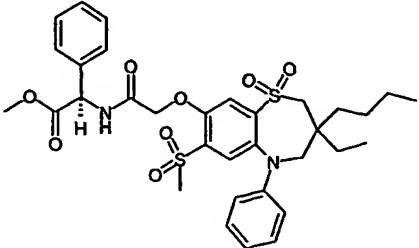
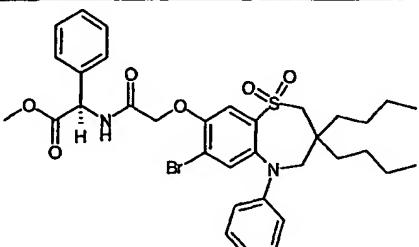
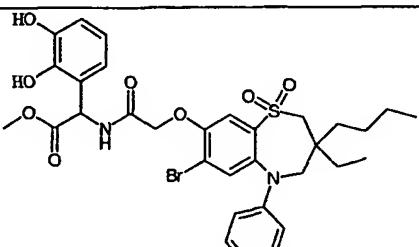
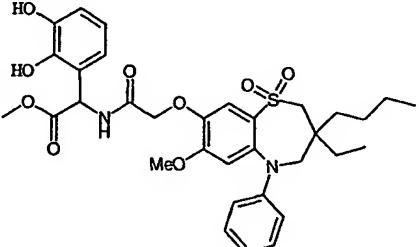
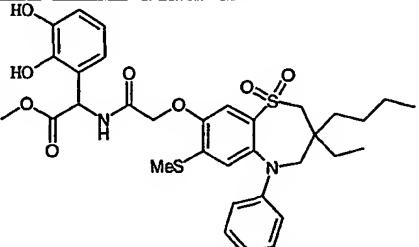
1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-((S)-1'-phenyl-1'-methoxycarbonylmethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 9; 50 mg, 0.098 mmol) was dissolved in DCM (2 ml). Methyl (2S)-amino(phenyl)acetate (19 mg, 0.12 mmol) and DIPEA (0.068 ml, 0.39 mmol) were added and the reaction was stirred for 2 minutes. TBTU (42 mg, 0.13 mmol) was added and the mixture was stirred for 1.5 hours at room temperature. The mixture was put on a pre-
10 packed ISOLUTE column and eluted with 10 ml DCM/EtOAc 8/2 to give the title compound 60 mg (93%). M/z 657.5.

Methods 47-62

The following compounds were synthesised by the procedure of Method 46 (except
15 that the reaction times were overnight) using the appropriate acid and amine (source not indicated where commercially available).

Meth	Compound	NMR or M/z	SM
47		609.4	Meth 10
48		625.4	Meth 17
49		685.3	Meth 14
50		609.4	Meth 10
51		637.4	Meth 15

52		657.4	Meth 21
53		685.3	Meth 14
54		0.73-0.95 (m, 6H), 0.98-1.78 (m, 8H), 3.12-3.28 (m, 2H), 3.6-4.0 (m, 5H), 4.60 (ABq, 2H), 5.79 (d, 1H), 6.0 (brs, 1H), 6.54 (dd, 1H), 6.83 (t, 1H), 6.95 (dd, 1H), 7.0-7.5 (m, 7H), 8.43 (d, NH), 9.32 (brs, 1H)	Meth 9 and Meth 74
55		0.75-0.9 (m, 6H), 1.0-1.78 (m, 8H), 3.10-3.26 (m, 2H), 3.63-3.87 (m, 8H), 4.56 (ABq, 2H), 5.76 (d, 1H), 5.99 (brs, 1H), 6.38 (s, 1H), 6.51 (dd, 1H), 6.81 (t, 1H), 6.93 (dd, 1H), 7.0-7.15 (m, 3H), 7.23-7.4 (m, 2H), 7.55 (s, 1H), 8.54 (d, NH), 9.45 (brs, 1H)	Meth 10 and Meth 74
56		0.76-0.87 (m, 6H), 1.0-1.8 (m, 8H), 2.23 (s, 3H), 3.1-3.25 (m, 2H), 3.6-3.95 (m, 5H), 4.61 (ABq, 2H), 5.79 (d, 1H), 6.0 (brs, 1H), 6.54 (dd, 1H), 6.65 (s, 1H), 6.83 (t, 1H), 6.92-7.1 (m, 4H), 7.23-7.4 (m, 3H), 8.37 (d, NH), 9.35 (brs, 1H)	Meth 17 and Meth 74

57		(CD ₃ OD) 0.76-0.85 (m, 6H), 1.02-1.3 (m, 8H), 1.36-1.56 (m, 4H), 2.16 (s, 3H), 3.24 (brs, 2H), 3.66-3.80 (m, 5H), 4.71 (ABq, 2H), 5.57 (s, 1H), 6.71 (s, 1H), 6.98 (t, 1H), 7.06-7.14 (m, 4H), 7.28 (brt, 2H), 7.37-7.45 (m, 3H)	Meth 22 and Meth 75
58		(CD ₃ OD) 0.76-0.85 (m, 6H), 1.02-1.28 (m, 8H), 1.36-1.56 (m, 4H), 1.96 (s, 3H), 3.24 (brs, 2H), 3.6-3.8 (m, 5H), 4.73 (ABq, 2H), 5.76 (s, 1H), 6.63 (s, 1H), 6.94-7.04 (m, 2H), 7.07-7.15 (m, 3H), 7.27 (t, 2H), 7.31 (s, 1H), 7.37 (d, 1H), 7.42 (s, 1H), 7.56 (d, 1H)	Meth 22
59		(CD ₃ OD) 0.80 (brt, 6H), 1.0-1.28 (m, 8H), 1.36-1.54 (m, 4H), 3.22 (brs, 2H), 3.61 (s, 3H), 3.69-3.8 (m, 5H), 4.62 (ABq, 2H), 5.56 (s, 1H), 6.49 (s, 1H), 6.99 (brt, 1H), 7.07-7.16 (m, 4H), 7.29 (brt, 2H), 7.37-7.43 (m, 2H), 7.52 (s, 1H)	Meth 6 and Meth 75
60		(CD ₃ OD) 0.75-0.84 (m, 6H), 1.0-1.29 (m, 8H), 1.35-1.54 (m, 4H), 3.20-3.23 (m, 5H), 3.65-3.8 (m, 5H), 4.64 (ABq, 2H), 5.74 (s, 1H), 6.34 (s, 1H), 6.95-7.04 (m, 2H), 7.09-7.15 (m, 3H), 7.24-7.31 (m, 3H), 7.37 (d, 1H), 7.50-7.54 (m, 2H)	Meth 6

61¹		0.74-0.83 (m, 6H), 0.98-1.7 (m, 8H), 3.18 (ABq, 2H), 3.60-3.90 (m, 5H), 4.59 (ABq, 2H), 5.67 (d, 1H), 7.0-7.2 (m, 4H), 7.2-7.55 (m, 8H), 7.91 (d, NH)	Meth 9
62		639.4	Meth 17 and Meth 76

¹ Eluent was DCM/EtOAc in a stepwise gradient from 100/0, 9/1, 8/2

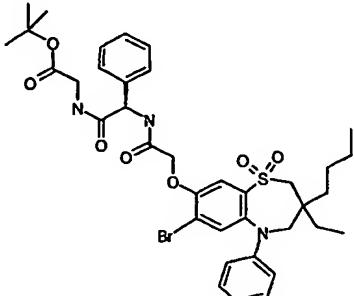
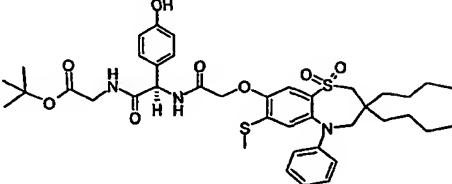
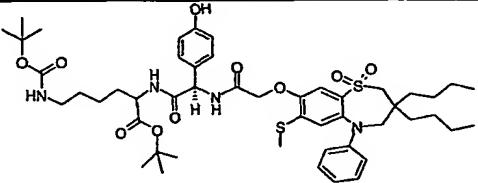
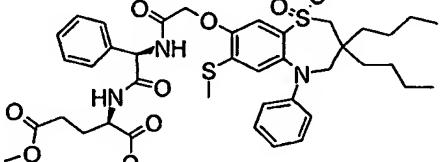
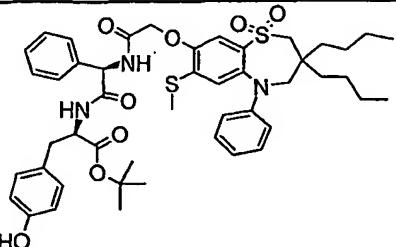
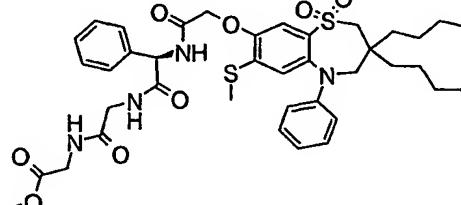
Method 63

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-[t-butoxycarbonylmethyl]carbamoyl)methyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 1; 110 mg, 0.17 mmol), glycine tert-butyl ester (30 mg, 0.23 mmol) and DIPEA (120 mg, 0.93 mmol) were dissolved in DCM (2 ml). The mixture was stirred for 5 mins at room temperature. TBTU (72 mg, 0.22 mmol) was added and the mixture was stirred for 1 h at room temperature. The solvent was evaporated at reduced pressure and the residue was placed on a column and the product was eluted with DCM/EtOAc (90:10) to give the title compound 122 mg (94%). NMR (300 MHz) 0.7-0.8 (m, 6H), 1.0-1.6 (m, 21H), 2.2 (s, 3H) 3.2 (s, 2H), 3.7-4.0 (m, 4H), 4.6 (ABq, 2H), 5.6 (d, 1H), 6.4 (t, 1H), 6.6 (s, 1H), 6.9-7.5 (m, 11H), 8.1 (d, 1H).

Methods 64-69

The following compounds were synthesised by the procedure of Method 63 using the appropriate acid and amine (source not indicated where commercially available).

Meth	Compound	NMR or M/z	SM
64		756.1	Ex 22
65		(CD ₃ OD) 0.75-0.85 (m, 6H), 1.1-1.3 (m, 8H), 1.4 (s, 9H), 1.45-1.55 (m, 4H), 2.15 (s, 3H), 3.25 (s, 2H), 3.75 (brs, 1H), 3.85 (s, 2H), 4.7 (ABq, 2H), 5.5 (s, 1H), 6.7 (s, 1H), 6.75-7.35 (m, 9H), 7.4 (s, 1H)	Ex 2
66		937.9 (M-H) ⁻	Ex 2
67		796.4	Ex 1
68			Ex 1
69			Ex 1

Method 70**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-[N-[(S)-1'-phenyl-1'-(diethoxyphosphoryl)methyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

The title compound was synthesised from 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 9) and diethyl (*S*)-amino(phenyl)methylphosphonate by the procedure of Example 33. NMR (600 MHz) 7.77-7.72 (1H, m), 7.47-7.42 (3H, m), 7.36-7.27 (5H, m), 7.14 (1H, s), 7.10-7.03 (2H, m), 5.55-5.48 (1H, m), 4.63-4.51 (2H, m), 4.14-4.02 (2H, m), 3.99-3.92 (1H, m), 3.81-3.60 (3H, m), 3.22-3.10 (2H, m), 1.65-1.25 (8H, m), 1.19-0.95 (6H, m), 0.78-0.73 (6H, m).

10

Method 71**4-Trifluoromethyl- α -methoxycarbonylbenzylamine**

4-Trifluoromethyl- α -carboxybenzylamine (1.4 g, 1.83 mmol) and thionylchloride were added to methanol (8 ml) and the mixture was refluxed for 2 h. The solvent was evaporated under reduced pressure. The residue was suspended in diethyl ether and the product was filtered off, washed with ether and dried to give the title compound 0.34 g (69%). NMR (300 MHz, DMSO-*d*₆) 3.3 (s, 1H), 5.45 (s, 1H), 7.7-7.9 (m, 4H), 9.25 (brs, 3H).

Method 72**20 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-[(R)-1'-phenyl-1'-(ethoxycarbonylmethyl)carbamoyl]methyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38; 52 mg, 0.082 mmol) and glycine ethyl ester hydrochloride (18 mg, 0.129 mmol) were dissolved in DCM (2 ml) and DIPEA (0.70 ml, 0.42 mmol) was added. After stirring at ambient temperature for 5 min TBTU (34 mg, 0.11 mmol) was added and the mixture was stirred for 2 hours. The solvent was evaporated and the residue was purified with flash chromatography (DCM:EtOAc 10:3) to give the title compound 50 mg (88%). NMR (500 MHz) 0.86 (m, 6H), 1.10-1.75 (m, 8H), 1.28 (m, 3H), 2.23 (s, 3H), 3.19 (q, 2H), 3.75 (m, 2H), 3.99-4.25 (m, 4H), 4.64 (q, 2H), 5.64 (m, 1H), 6.35 (brs, 1H), 6.69 (s, 1H), 7.03 (t, 1H), 7.09 (d, 1H), 7.29-7.52 (m, 7H), 8.10 (d, 1H).

Method 73

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N-(1"-methoxycarbonyl-1"-phenylmethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

5 The title compound was synthesised by the procedure of Method 72 using 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-((R)-1'-phenyl-1'-carboxymethyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 38) and methyl (2R)-amino(phenyl)acetate hydrochloride.

10 **Method 74**

1-(1'-Methoxycarbonyl-1'-aminomethyl)-2,3-dihydroxyphenyl hydrochloride salt

1-(1'-Carboxy-1'-aminomethyl)-2,3-dihydroxyphenyl (40 g, 0.218 m mol) was mixed with methanol (230 ml). HCl gas was bubbled through. The mixture was refluxed for 2 hours. The solvent was evaporated under reduced pressure. The product was crystallised from 15 methanol/EtOAc/diethyl ether to yield 35.5 g (70 %) of the title product. NMR (600 MHz, CD₃OD) 3.76 (s, 3H), 5.19 (s, 1H), 6.68-6-75 (m, 2H), 6.85 (dd, 1H)

Method 75

(R)-1-(1'-Methoxycarbonyl-1'-aminomethyl)-4-fluorophenyl hydrochloride salt

20 (2R)-amino(4-fluorophenyl)acetic acid (570 mg, 2.77 mmol) was dissolved in methanol (5 ml) and cooled in an ice-bath. Thionyl chloride (2 ml) was added dropwise and temperature was allowed to reach room temperature. After 5 hours the mixture was evaporated under reduced pressure. The procedure was repeated and the reaction was stirred overnight. The mixture was evaporated under reduced pressure to give the title product in a 25 quantitative yield. NMR (500 MHz, CD₃OD) 3.84 (s, 3H), 5.26 (s, 1H), 7.26 (t, 2H), 7.53 (dd, 2H).

Methods 76-77

The following compounds were synthesised by the procedure of Method 75 using the 30 appropriate acid and amine (source not indicated where commercially available).

Meth	Compound	NMR	SM
76	(S)- α -Methylamino- α -methoxycarbonylbenzyl	(CD ₃ OD) 2.63 (s, 3H), 3.81 (s, 3H), 5.15 (s, 1H), 7.45-7.55 (m, 5H)	(S)- α -Methylamino- α -carboxybenzyl
77 ¹	α -Methoxycarbonyl-N-methylbenzylamine hydrochloride	(D ₂ O) 2.65 (s, 3H), 3.81 (s, 3H), 5.15 (s, 1H), 7.45-7.48 (m, 2H), 7.52-7.59 (m, 3H)	(methylamino)(phenyl)acetic acid

¹ Total reaction time 5 days

Method 78

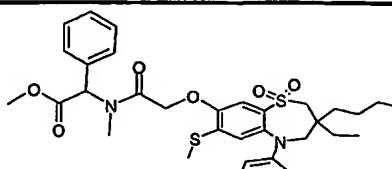
5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-[(R)- α -(*t*-butoxycarbonyl)-4-hydroxybenzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

tert-Butyl (2R)-amino(4-hydroxyphenyl)acetate (104 mg, 0.47 mmol) and 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 17; 185 mg, 0.39 mmol) were dissolved in DCM (5 ml) and lutidine 10 (0.09 ml, 0.77 mmol) was added. After stirring at room temperature for 5 min o-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (208 mg, 0.55 mmol) was added and the reaction mixture was stirred at room temperature for 2 hours.

Purification with flash chromatography (DCM:EtOAc 10:1→5:1) gave the title compound (175 mg, 66 %). NMR (300 MHz) 0.81 (m, 6H), 1.05-1.65 (m, 8H), 1.42 (s, 9H), 2.21 (s, 3H), 15 3.17 (ABq, 2H), 3.74 (m, 2H), 4.60 (ABq, 2H), 5.22 (brs, 1H), 5.49 (d, 1H), 6.67 (s, 1H), 6.79 (m, 2H), 7.00 (t, 1H), 7.07 (d, 2H), 7.23-7.30 (m, 3H), 7.40 (s, 1H), 7.82 (brd, 1H).

Method 79

The following compounds were synthesised by the procedure of Method 78 using the 20 appropriate starting material.

Meth	Compound	M/z	SM
79		639.3	Meth 17 and Meth 77

Method 80**2-{[(2R)-2-Amino-2-(4-hydroxyphenyl)ethanoyl]amino}ethanesulphonic acid**

N-Boc-4-hydroxyphenylglycine (1.00 g, 3.21 mmol) was dissolved in DMF (5 ml) and tetrabutylammonium taurine (2.36 g, 6.42 mmol) was added together with additionally 5 ml 5 DMF. The resulting suspension was cooled on ice and TBTU (1.24 g, 3.85 mmol) was added. The ice bath was removed after 30 min and the mixture was stirred for 2 hours before it was filtered and concentrated. TFA in DCM (20%, 20 ml) was added and the reaction mixture was stirred overnight. Ethanol (20 ml) was added and the solvents evaporated. The crude product was refluxed in ethanol (100 ml) for 1 hour. Filtration yielded the pure title compound as a 10 white solid, 626 mg (71%). NMR (DMSO-d₆) 2.4-2.6 (m, 2H), 3.2-3.4 (m, 2H), 4.79 (s, 1H), 6.78 (d, 2H), 7.23 (d, 2H), 8.22 (t, 1H), 8.4 (brs, 3H), 9.7 (s, 1H).

Method 81**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-carboxymethylthio-8-methoxy-2,3,4,5-tetrahydro-1,5-****benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (113 mg, 0.24 mmol), Cs₂CO₃ (170 mg, 0.52 mmol) and ethyl thioglycolate (0.060 ml, 0.54 mmol) in DMF (4.0 ml) were subjected to microwave irradiation in a Smith Synthesiser at 80°C for 3 min and then at 90°C for 8 min. The reaction mixture was diluted 20 with water (250 ml) and extracted with DCM (5 x 10 ml) and collected organic layers were dried (MgSO₄), concentrated and purified on a short column (petroleum ether : EtOAc 4:1 → 2:1). The resulting product was dissolved in THF (2 ml) and water (2 ml) and NaOH (aq., 0.5 ml, 1 M) was added and the reaction mixture was stirred at room temperature for 2 hours. The reaction was quenched with HCl (1 M) and the reaction mixture was diluted with water (10 ml) and extracted with DCM (3 x 3 ml). Purification with preparative HPLC yielded the title 25 compound (58 mg, 59 %). NMR (300 MHz, CD₃OD) 0.81 (m, 6H), 1.00-1.70 (m, 8H), 3.21 (m, 2H), 3.42 (m, 2H), 3.71 (m, 2H), 3.92 (s, 3H), 6.88 (m, 2H), 7.02 (m, 2H), 7.23 (t, 2H), 7.40 (s, 1H).

Method 82**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-ethoxycarbonylmethylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-carboxymethoxy-2,3,4,5-tetrahydro-
5 1,5-benzothiazepine (Method 9; 50 mg, 0.098 mmol) and Cs₂CO₃ (51 mg, 0.15 mmol) were
added to DMF (2.0 ml) and ethyl thioglycolate (0.02 ml, 0.15 mmol) was added. The reaction
mixture was subjected to microwave irradiation in a Smith Synthesiser at 150°C for 5 min.
The reaction mixture was diluted with water (100 ml), made acidic with HCl (1 M), extracted
with DCM (3 x 10 ml) and the collected organic layers were dried (MgSO₄) to give the crude
10 title compound (54 mg). M/z 550.2.

Method 83 and Method 84**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Enantiomer 1); and**

15 **1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Enantiomer 2)**

The two enantiomers of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-
2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 96/16051) were obtained by separation of the
corresponding racemic mixture using preparative HPLC. The column used was a Chiraldak
20 AD (20x250 mm i.d., 10 \square m) and the mobile phase was a heptane / IPA mixture in ratio
90/10. The injected racemate (17.3 mg in IPA (1 ml)) was eluted with a flow of 10 ml/min
and the chromatogram was followed with UV-detection at 285nm. Totally 260 mg racemate
was separated yielding 121 mg of the first eluting enantiomer (Enantiomer 1) and 115 mg of
the second eluting enantiomer (Enantiomer 2). Total yield 91%. Each of the two enantiomers
25 was obtained in 99.4% e.e.

Method 85**(R)-N-Benzylloxycarbonyl- α -[N'-(t-butoxycarbonylmethyl)carbamoyl]benzylamine**

(R)-N-Benzylloxycarbonyl- α -carboxybenzylamine (10 g, 35.0 mmol) and t-
30 butylglycine hydrochloride (6.3 g, 37.4 mmol) was dissolved in DCM (200 ml) with 2,6-
lutidine (8.2 ml, 70.4 mmol). After stirring 5 min at 0°C TBTU (12.4 g, 38.6 mmol) was
added and stirring was continued 1 hour 30 min at 0°C and 3 hours 45 min at room
temperature. The reaction mixture was washed with water (2 x 100 ml), dried (MgSO₄) and

purified with flash chromatography (DCM:EtOAc 7:1→5:1) to give the title compound (13 g, 94 %). NMR (500 MHz) 1.45 (s, 9H), 3.84 (d, 1H), 4.00 (dd, 1H), 5.10 (m, 2H), 5.28 (brs, 1H), 6.13 (brs, 1H), 6.23 (brs, 1H), 7.30-7.44 (m, 10H).

5 Method 86

(R)- α -[N-(*t*-Butoxycarbonylmethyl)carbamoyl]benzylamine

(R)-N-Benzylloxycarbonyl- α -[N'-(*t*-butoxycarbonylmethyl)carbamoyl]benzylamine (Method 85; 12.8 g, 32.2 mmol) was dissolved in EtOH (99%, 200 ml) and toluene (50 ml). Pd/C (10%, 0.65 g) was added and hydrogenation was performed at atmospheric pressure for 10 5 hours 30 min at room temperature. The reaction mixture was filtered through diatomaceous earth and the solvents were evaporated to give the title compound (8.4 g, 99 %). NMR (600 MHz) 1.45 (s, 9H), 3.93 (m, 2H), 4.54 (s, 1H), 7.31-7.42 (m, 5H), 7.51 (brs, 1H).

Method 87

15 1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(α-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 22; 50 mg, 0.099 mmol) was dissolved in DCM (2 ml). (S)-Phenylglycine methyl ester hydrochloride (24.8 mg, 0.123 mmol) and diisopropyl ethyl amine 20 (70 □l, 0.401 mmol) were added. The mixture was stirred for 15 min and then TBTU (38 mg, 0.118 mmol) was added. The reaction was completed after 1.5h (LC/MS). The crude product was purified by flash chromatography using chloroform/EtOAc (8/2) as the eluent (88.6%; 55.2 mg, 0.064 mmol). M/z 653.

25 Method 88

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)- α -[N'-(methoxycarbonylmethyl)carbamoyl]benzyl]carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-[N-(S)-(α-carboxybenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Example 88; 25 mg, 0.039 mmol) and glycine methyl ester (7.5 mg, 0.059 mmol) were dissolved in DCM (2 ml). Diisopropyl ethyl amine (27 □l, 0.158 mmol) and TBTU (15 mg, 0.047 mmol) were added successively and the mixture was stirred for 2 hours at ambient temperature. The crude

product was purified by flash chromatography using DCM/EtOAc (8/2) as eluent 79% yield (22 mg). M/z 710.

Method 89

5 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-(2-carboxyethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

Sodium hydroxide (38 mg, 0.95 mmol) was dissolved in ethanol (2.5 ml) and then 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 96/16051; 200 mg, 0.443 mmol) was added. After stirring at room temperature for 5 min, 3-bromopropionic acid (68 mg, 0.443 mmol) was added and the reaction mixture was refluxed for 20 hours. Acetic acid was added. The solvent was evaporated under reduced pressure and the residue was extracted with EtOAc/water. The organic layer was separated, washed with water, dried and evaporated under reduced pressure. The crude product was purified by column chromatography using DCM/MeOH (100:5) as eluent to give the title compound 89 mg (38%). NMR (CD_3OD) 0.75-0.83 (m, 6H), 1.0-1.25 (m, 4H), 1.38-1.65 (m, 4H), 2.82 (m, 2H), 3.26 (s, 2H), 3.50-3.90 (m, 2H), 4.33 (t, 2H), 6.99 (t, 1H), 7.07-7.13 (m, 3H), 7.28 (t, 2H), 7.53 (s, 1H).

Method 90

20 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-(2-{N-[{(R)-\alpha-(t-butoxycarbonyl)benzyl]carbamoyl}ethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-bromo-8-(2-carboxyethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 89; 70 mg, 0.134 mmol) and (R)-\alpha-(t-butoxycarbonyl)benzylamine (J.Amer.Chem.Soc.; EN; 117; 44; 1995; 10879-10888; 35 mg, 0.169 mmol) in DCM (2.5 ml) was added 2,6-lutidine (29 mg, 0.268 mmol) and TBTU (56 mg, 0.174 mmol). The reaction mixture was stirred at room temperature for 2.5 hours, then diluted with DCM. The solution was washed with NaHCO_3 (aq, sat), water, dried and the solvent was evaporated under reduced pressure. The residue was suspended in ether/petroleum ether and the crystals filtered to give the title compound 85 mg (89%). NMR (500 MHz) 0.79-0.86 (m, 6H), 1.04-1.28 (m, 4H), 1.35-1.56 (m, 11H), 1.60-1.77 (m, 2H), 2.82 (t, 2H), 3.13-3.25 (m, 2H), 3.72 (brs, 2H), 4.35-4.44 (m, 2H), 5.54 (d, 1H), 6.95 (d, 1H), 7.04 (t, 1H), 7.08 (d, 2H), 7.15 (s, 1H), 7.29-7.43 (m, 6H), 7.52 (s, 1H).

Methods 91-94

The following compounds were synthesised by the procedure of Example 104 using the appropriate starting material (source of amine indicated where not commercially available).

Meth	Compound	NMR (500 MHz) and m/z	SM
91		0.77-0.86 (m, 6H), 1.03-1.62 (m, 21H), 2.21 (s, 3H), 2.32 (dd, 1H), 2.54 (dd, 1H), 3.14 (s, 2H), 3.74 (brs, 2H), 4.48-4.53 (m, 1H), 4.60 (dd, 2H), 5.57 (d, 1H), 6.33 (d, 1H), 6.67 (s, 1H), 7.01 (t, 1H), 7.09 (d, 2H), 7.17-7.40 (m, 21H), 7.50 (d, 2H), 8.10 (d, 1H); m/z 1040.83	Ex 1; 1
92		0.78-0.86 (m, 6H), 1.05-1.27 (m, 8H), 1.36-1.58 (m, 13H), 1.78 (s, 3H), 2.23 (s, 3H), 2.77-2.92 (m, 2H), 3.19 (s, 2H), 7.75 (brs, 2H), 4.64 (dd, 2H), 4.72-4.77 (m, 1H), 6.68 (s, 1H), 6.81 (d, 1H), 7.01 (t, 1H), 7.09 (d, 2H), 7.27-7.42 (m, 6H), 7.50 (d, 2H), 8.16 (d, 1H); m/z 812.23	Ex 1; 2
93		0.74-0.81 (m, 6H), 1.0-1.22 (m, 8H), 1.29-1.62 (m, 13H), 2.13 (s, 3H), 2.50-2.64 (m, 2H), 3.14 (s, 2H), 3.69 (brs, 2H), 4.42-4.48 (m, 1H), 4.58 (dd, 2H), 5.45 (d, 1H), 6.13 (d, 1H), 6.62 (s, 1H), 6.96 (t, 1H), 7.04 (d, 2H), 7.17-7.21 (m, 3H), 7.23-7.37 (m, 18H), 7.41 (d, 2H), 8.0 (d, 1H)	Ex 1; Meth 113

94		0.81-0.87 (m, 6H), 1.06-1.29 (m, 8H), 1.39-1.61 (m, 4H), 1.78 (brs, 2H), 1.94 (s, 3H), 2.07-2.17 (m, 1H), 2.20- 2.27 (m, 4H), 3.31 (s, 2H), 3.77 (brs, 2H), 3.80 (s, 3H), 4.65 (dd, 2H), 4.76- 4.82 (m, 1H), 5.65-5.70 (m, 1H), 6.69 (s, 1H), 7.04 (t, 1H), 7.12 (d, 2H), 7.29-7.44 (m, 7H), 7.52 (d, 2H), 8.16 (d, 1H)	Ex 1
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¹ *t*-butyl L-(S-trityl)cysteinate hydrochloride: Org. Pre. Proced. Int.; 1999, 31:571-572

² S-methyl-L-cysteine tert-butyl ester: Pestic. Sci.; EN; 45; 4; 1995; 357-362

Method 95

5 3,3-Dibutyl-4-oxo-5-(4-chlorophenyl)-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

A mixture of 3,3-dibutyl-4-oxo-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (WO 95/04534; 1.0 g, 2.5 mmol), 4-bromochlorobenzene (4.78 g, 24.98 mmol), copper bromide (36 mg, 0.25 mmol) and potassium carbonate (0.35 g, 2.5 mmol) was refluxed for 20 hours. The reaction mixture was loaded onto a column and the product eluted with 5% EtOAc/petroleum ether (0.8 g, 63% yield). NMR (500 MHz) 0.86-0.92 (m, 6H), 1.16-1.35 (m, 8H), 1.45-1.65 (m, 4H), 3.16 (s, 2H), 3.96 (s, 3H), 7.06-7.10 (m, 2H), 7.19 (s, 1H), 7.29 (s, 1H), 7.33-7.38 (m, 2H). M/z 511.

15 Method 96

1,1-Dioxo-3,3-dibutyl-4-oxo-5-(4-chlorophenyl)-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a mixture of 3,3-dibutyl-4-oxo-5-(4-chlorophenyl)-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 95; 0.67 g, 1.304 mmol), DCM (34 ml), water (34 ml) and potassium carbonate (0.554 g, 4.0 mmol) was added at 0 °C m-chloroperoxybenzoic acid (0.78 g, 3.2 mmol) in one portion. The reaction mixture was stirred at 0 °C for 10 h and then at room temperature for 14 hours. DCM (100 ml) and NaHCO₃ (aq, sat; 150 ml) were added. The organic layer was separated, washed with brine, dried and evaporated under reduced pressure to give the title compound 0.68 g (96%). NMR (600 MHz) 0.7-0.92 (m, 6H),

1.0-1.60 (m, 10H), 1.70-1.92 (m, 2H), 2.30-3.7 (m, 2H), 3.99 (s, 3H), 7.16-7.20 (m, 2H), 7.24 (s, 1H), 7.34-7.37 (m, 2H), 7.44 (s, 1H); m/z 543.

Method 97

5 1,1-Dioxo-3,3-dibutyl-4-oxo-5-(4-chlorophenyl)-7-methylthio-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

Sodium methanethiolate (0.43 g, 6.08 mmol) was added to a solution of 1,1-dioxo-3,3-dibutyl-4-oxo-5-(4-chlorophenyl)-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 96; 0.66 g, 1.22 mmol) in anhydrous DMF (11 ml) under nitrogen.
10 The reaction mixture was stirred at room temperature for 72 hours. The solvent was evaporated under reduced pressure and the residue was extracted with trichloromethane/water. The organic layer was separated, washed with brine, dried and evaporated under reduced pressure. The crude product was purified by column chromatography using DCM as eluent to give the title compound 0.6 g (96%). NMR (500 MHz) 0.80-1.0 (m, 6H), 1.10-1.6 (m, 10H),
15 1.70-2.0 (m, 2H), 2.28 (s, 3H), 3.37-3.70 (m, 2H), 4.04 (s, 3H), 6.65 (s, 1H), 7.25-7.30 (m, 2H), 7.35-7.42 (m, 3H); m/z 510.4.

Method 98

1,1-Dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3,3-dibutyl-4-oxo-5-(4-chlorophenyl)-7-methylthio-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 97; 0.41 g, 0.79 mmol) in anhydrous ether (15 ml) was added LiAlH₄ (0.15 g, 3.97 mmol) under nitrogen. The reaction mixture was stirred at room temperature for 2.5 hours. The reaction flask was cooled to 0°C
25 and the excess LiAlH₄ was quenched by adding water (0.3 ml) and 2 M aqueous NaOH (0.3 ml). The mixture was filtered and the filtrate was dried and evaporated under reduced pressure. The crude product was purified by column chromatography using DCM as eluent to give the title compound 0.265 g (68%). NMR (300 MHz) 0.8-0.90 (m, 6H), 1.0-1.47 (m, 12H), 2.33 (s, 3H), 3.17 (s, 2H), 3.70 (s, 2H), 3.93 (s, 3H), 7.03-7.08 (m, 3H), 7.23-7.32 (m, 3H); m/z 496.

Method 99**1,1-Dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-methoxy-
5 2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 98; 0.26 g 0.52 mmol) in anhydrous DCM
(10 ml) was added boron tribromide (2.63 g, 10.48 mmol) under nitrogen. The reaction
mixture was stirred at room temperature for 2.5 h. The reaction flask was cooled to 0°C, water
(20 ml) and hydrazine monohydrate (0.5 ml) was added. The organic layer was separated,
dried and evaporated under reduced pressure. The crude product was purified by column
10 chromatography using DCM/EtOAc (100:5 and 100:10) as eluent to give the title compound
0.20 g (80%). NMR (500 MHz) 0.85 (t, 6H), 1.03-1.28 (m, 8H), 1.35-1.46 (m, 4H), 2.39 (s,
3H), 3.21 (s, 2H), 3.73 (s, 2H), 7.04 (d, 2H), 7.29-7.34 (m, 3H), 7.44 (s, 1H); m/z 482.

Method 100**1,1-Dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

Ethyl bromoacetate (0.101 g, 0.604 mmol) was added to a mixture of 1,1-dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 99; 0.194 g, 0.402 mmol), anhydrous Na₂CO₃ (0.192 g, 1.81 mmol) and
20 tetrabutylammonium bromide in MeCN (5 ml). The reaction mixture was refluxed for 3.5 hours. The solvent was evaporated under reduced pressure and the residue was extracted with DCM/water. The organic layer was separated, dried and evaporated under reduced pressure. The crude product was purified by column chromatography using DCM/EtOAc (100:5 and 100:10) as eluent to give the title compound 0.197 g (86%). NMR (300 MHz) 0.80-0.89 (m, 6H), 1.0-1.45 (m, 15H), 2.34 (s, 3H), 3.16 (s, 2H), 3.68 (s, 2H), 4.30 (q, 2H), 4.71 (s, 2H),
25 7.05-7.11 (m, 3H), 7.19 (s, 1H), 7.29-7.35 (m, 2H).

Method 101**1,1-Dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a solution of 1,1-dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 100; 0.195 g, 0.343 mmol) in ethanol (8 ml) was added NaOH (1.03 mmol in 0.5 ml water). The reaction mixture

was stirred at room temperature for 70 min and then quenched by adding acetic acid (0.3 ml). The solvent was evaporated under reduced pressure and the residue was extracted with DCM/water. The organic layer was separated, washed with brine, dried and evaporated under reduced pressure to give the title compound 0.169 g (91%). NMR (500 MHz, CD₃OD) 0.86 (t, 5 6H), 1.11-1.28 (m, 8H), 1.37-1.44 (m, 4H), 2.33 (s, 3H), 3.25 (s, 2H), 3.55 (s, 2H), 4.73 (s, 2H), 7.10-7.15 (m, 3H), 7.26 (s, 1H), 7.28-7.32 (m, 2H).

Method 102

1,1-Dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-[N-[(R)-α-[N'-(t-butoxycarbonylmethyl)carbamoyl]benzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a solution of 1,1-dioxo-3,3-dibutyl-5-(4-chlorophenyl)-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 101; 100 mg, 0.185 mmol) and (R)-α-[N-(*t*-butoxycarbonylmethyl)carbamoyl]benzylamine (Method 86; 56 mg, 0.213 mmol) in DCM (4 ml) was added 2,6-lutidine (40 mg, 0.37 mmol) and TBTU (89 mg, 0.28 mmol). The reaction mixture was stirred at room temperature for 2 hours and then EtOAc was added and the solution washed with water. The organic layer was separated, dried and evaporated under reduced pressure. The crude product was purified by column chromatography using DCM/MeOH (100:3) as eluent to give the title compound 0.129 g (89%). NMR (600 MHz) 0.78-82 (m, 6H), 1.01-1.23 (m, 8H), 1.30-1.42 (m, 13H), 2.32 (s, 3H), 3.10-3.16 (m, 2H), 3.62-3.68 (m, 2H), 3.81-3.87 (m, 1H), 3.95-4.03 (m, 1H), 4.52 (dd, 2H), 5.57 (d, 1H), 6.27 (t, 1H), 7.01-7.07 (m, 3H), 7.20-7.43 (m, 8H), 8.02 (d, 1H).

Method 103

3,3-Dibutyl-4-oxo-5-(4-nitrophenyl)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To 3,3-dibutyl-4-oxo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (synthesised by the procedure of WO9616051 for the corresponding 3-butyl-3-ethyl analogue; 2.9 g, 9.0 mmol) was added p-nitrophenylbromide (24 g, 119 mmol), K₂CO₃ (1.6 g, 12 mmol) and CuI (180 mg, 0.95 mmol). The reaction mixture was heated to 200°C overnight. Then it was allowed to cool down to room temperature and the resulting solid was purified by chromatography using DCM as eluent. The fractions containing the product were concentrated under reduced pressure and EtOH (95%) was added and the insoluble p-nitrophenylbromide was then filtered off. The residue was purified again by flash

chromatography using DCM as eluent. The product was still not pure so the residue was then purified by flash chromatography using EtOAc:heptane, 1:9 as eluent to give the title compound 2.57 g (64%). NMR (600 MHz) 0.77-0.87 (m, 6H), 1.12-1.31 (m, 8H), 1.4-1.6 (m, 4H), 3.09 (brs, 2H), 3.79 (s, 3H), 6.72-6.83 (m, 2H), 7.18-7.27 (m, 3H), 8.3 (d, 2H).

5

Method 104

1,1-Dioxo-3,3-dibutyl-4-oxo-5-(4-nitrophenyl)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To 3,3-dibutyl-4-oxo-5-(4-nitrophenyl)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 103; 2.57 g, 5.8 mmol) was added DCM (130 ml), water (130 ml) and K₂CO₃ (2.44 g, 17.6 mmol). The reaction mixture was cooled down to 0°C and m-chloroperoxybenzoic acid (3.42 g, 13.9 mmol) was added in one portion. The reaction was allowed to complete overnight with the temperature slowly rising to room temperature. Then NaHCO₃aq (sat) was added and the two layers were separated. The water layer was then extracted three times with DCM. The combined organic layers was dried, filtered and evaporated under reduced pressure. The product was purified by flash chromatography using DCM as eluent to give the title compound 2.4 g (87%). M/z 475.4.

Method 105

1,1-Dioxo-3,3-dibutyl-5-(4-aminophenyl)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To LiAlH₄ (5.76 g, 151 mmol) were added THF (200 ml). The reaction mixture was cooled to 0 °C. and H₂SO₄ (4.06 ml, 76 mmol) were added slowly with a syringe. After the addition was completed the reaction was stirred for 10 minutes. Then 1,1-dioxo-3,3-dibutyl-4-oxo-5-(4-nitrophenyl)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 104; 2.57 g, 5.06 mmol) dissolved in THF (50 ml) was added at 0°C. After vigorous stirring for 1hour the cooling bath was removed and the reaction was heated to 40°C overnight. Then Na₂SO₄.10H₂O (3-4 teaspoons), water (8 ml), NaOH (15%, aq) (8 ml), water (25 ml) and MeOH (30 ml) were added in that order. The precipitate was removed by filtration and rinsed with DCM/MeOH. The solvent was dried, filtered and concentrated under reduced pressure.

The residue was purified by flash chromatography using DCM:EtOAc, 9:1 then 3:1 as eluent to give the title compound 0.6 g (27%). M/z 431.3.

Method 106**1,1-Dioxo-3,3-dibutyl-5-(4-aminophenyl)-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3,3-dibutyl-5-(4-aminophenyl)-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 105; 918 mg, 2.13 mmol) was dissolved in DMF (dry, 20 ml).

- 5 Sodium thiomethoxide (810 mg, 11.6 mmol) was added. The reaction mixture was treated at 100-120°C for four hours then room temperature overnight. Acetic acid (3 ml) was added and the mixture was flushed with nitrogen (g) and the gas was lead through a flask containing sodium hypochlorite in order to destroy the methyl mercaptan formed. Water was added and the water layer was extracted two times with EtOAc. The combined organic layer was washed
- 10 with brine, dried, filtered and evaporated under reduced pressure. The mixture contained DMF so toluene and brine was added (everything didn't dissolve). The water layer was extracted two times with toluene. The combined organic layers were washed once with brine. The separation funnel was washed with EtOAc in order to dissolve everything. The toluene and EtOAc solutions were combined, dried, filtered and evaporated under reduced pressure.
- 15 The residue was purified by flash chromatography using DCM:EtOAc, 7:3 as eluent to give the title compound 0.6 g (27%). M/z 417.4.

Method 107**1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

- 20 1,1-Dioxo-3,3-dibutyl-5-(4-aminophenyl)-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 106; 600 mg, 1.44 mmol) was dissolved in THF (10 ml). Di-t-butyl dicarbonate (314 mg, 1.44 mmol) was added and the mixture was stirred at 60°C for two hours and room temperature for 3 days. The solvent was evaporated under reduced pressure.
- 25 EtOAc was added and the organic layer was washed once with KHSO₄-solution (0.3M, aq) and once with brine, dried, filtered and evaporated under reduced pressure. The residue was purified by flash chromatography using DCM:EtOAc, 9:1 as eluent to give the title compound 0.597 g (80%). M/z 517.3.

Method 108

1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 107; 597 mg, 1.16 mmol) was dissolved in MeCN (20 ml), K₂CO₃ (480 mg, 3.5 mmol), tetrabutylammoniumbromide (54 mg, 0.17 mmol) and ethyl bromoacetate (167 µl, 1.5 mmol) was added. The mixture was heated to 60°C overnight. The solvent was evaporated under reduced pressure. EtOAc and water was added and the water layer was extracted two times with EtOAc. The combined organic layer was washed once with brine, dried, filtered and evaporated under reduced pressure to give the title compound 0.617 g (89%). M/z 603.3.

Method 109

1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

15 1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 108; 607 mg, 1.0 mmol) was dissolved in THF (6 ml), H₂O (6 ml) and LiOH (127 mg, 3.02 mmol, monohydrate) was added. The mixture was stirred for 1 hour. The mixture was poured into 20 water and the solution was acidified using HCl-solution (aq, 1M). The water layer was extracted two times with EtOAc. The combined organic layer was wash once with brine, dried, filtered and evaporated under reduced pressure to give the title compound 0.571 g (99%). M/z 575.4.

25 **Method 110**

1,1-Dioxo-3,3-dibutyl-5-(4-aminophenyl)-8-[N-(α-(R)-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-(4-t-butoxycarbonylaminophenyl)-8-[N-(α-(R)-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 30 45; 562 mg, 0.78 mmol) was dissolved in DCM (18 ml). TFA (4 ml) was added and the reactions mixture was stirred for 3 hours. The solvent was evaporated under reduced pressure. The residue was partitioned between EtOAc and NaOH solution (1M, aq). The aqueous phase was extracted one more time with EtOAc. The combined organic layer was wash with brine,

dried, filtered and evaporated under reduced pressure to give the title compound 440 mg (91%). M/z 622.5.

Method 111

5 1,1-Dioxo-3,3-dibutyl-5-[4-(N'-*t*-butylureido)phenyl]-8-[N-(α -(R)-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

1,1-Dioxo-3,3-dibutyl-5-(4-aminophenyl)-8-[N-(α -(R)-methoxycarbonylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 110; 40 mg, 0.064 mmole) was dissolved in DMF (1 ml). *t*-Butyl isocyanate (8.3 μ l, 0.071 mmol) was added. The 10 reaction mixture was stirred at 60-80°C overnight. Tert-butyl isocyanate (20 μ l, 0.171 mmol) was added. The reaction mixture was stirred at 60-80°C for 2 days and then room temperature for a few days. The solvent was evaporated under reduced pressure. The product was purified by preparative HPLC using an MeCN/ammonium acetate buffer gradient (5/95 to 100/0) as eluent to give the title product, 30 mg (65%). M/z 721.6.

15

Method 112

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-[N-(α -methoxycarbonylmethylbenzyl)carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine

The title compound was synthesized from 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 17) and methyl 3-amino-3-phenylpropanoate (Helv.Chim.Acta; EN; 83; 6; 2000; 1256 – 1267) by the procedure of Example 56. M/z 639.4.

Method 113

25 *t*-Butyl D-(S-trityl)cysteinate hydrochloride

To a vigorously stirred suspension of S-trityl-D-cysteine (2.0 g, 5.5 mmol) in *t*-butyl acetate (35 ml), 70% HClO₄ (1.6 ml) was added dropwise. The reaction mixture was stirred at room temperature for 70 min and EtOAc (50 ml) and NaHCO₃ (aq, sat) to pH 8.0 were added. The precipitate, unreacted S-trityl-D-cysteine was filtered off. The organic layer was separated, washed with 0.5 M HCl (2 x 75 ml) and brine, dried and evaporated to give the title compound 2.02 g (81%). NMR (500 MHz): 1.43 (s, 9H), 2.83-2.95 (m, 2H), 3.41-3.48 (m, 1H), 7.21-7.37 (m, 9H), 7.46 (d, 6H).

Method 114**1,1-Dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-ethoxycarbonyl-2,3,4,5-tetrahydro-1,5-benzothiazepine**

To a suspension of 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 26; 12.85 g, 28.71 mmol) in MeCN (150 ml) was added ethyl bromoacetate (3.85 ml, 34.6 mmol), tetrabutylammonium bromide (0.925 g, 2.869 mmol) and sodium carbonate (12.85 g, 121.2 mmol). The mixture was heated under reflux for 5 hours. The solvent was removed under reduced pressure and the residue was partitioned between DCM and 0.5 M HCl. The organic layer was washed with brine, dried (MgSO₄) and concentrated. Chromatography using DCM/EtOAc (9:1) as eluent gave the desired product (15.45 g) as a tan oil. NMR 0.70-0.85 (m, 6H), 1.00-1.55 (m, 15H), 2.15 (s, 3H), 3.10 (s, 2H), 3.70 (bs, 2H), 4.25 (q, 2H), 4.70 (s, 2H), 6.65 (s, 1H), 6.90-7.30 (m, 6H).

Method 115**15 1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 116; 0.48 g, 1.04 mmol) was dissolved in ethanol (10 ml). NaOH (0.30 g, 7.5 mmol) was added and the mixture was refluxed for 30 min. Acetic acid (1 ml) was added. The solvent was evaporated under reduced pressure and the residue was extracted with DCM/water. The DCM layer was separated, dried and evaporated. 0.44 g (97%) of the title compound was obtained. NMR (300 MHz) 0.7-0.8 (m, 6H), 1.0-1.6 (m, 8H), 3.1-3.3 (m, 2H), 3.5-3.8 (m, 2H), 4.6 (s, 3H), 6.8-7.3 (m, 7H), 7.5 (s, 1H).

25 Method 116**1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-8-ethoxycarbonylmethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine**

1,1-Dioxo-3-butyl-3-ethyl-5-phenyl-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepin (WO9616051; 0.40 g, 1.07 mmol), ethyl bromoacetate (0.23 g, 1.38 mmol), sodium carbonate (0.50 g, 4.7 mmol) and tetrabutylammonium bromide (30 mg, 0.093 mmol) were added to MeCN (10 ml). The mixture was refluxed for 18 hours and then evaporated under reduced pressure. The residue was extracted with DCM/water. The DCM layer was separated and evaporated under reduced pressure. The residue was purified by column chromatography. The

product was eluted with DCM/EtOAc (90:10). 0.480 g (97%) of the title compound was obtained. NMR (300 MHz) 0.7-0.85 (m, 6H), 1.0-1.7 (m, 11H), 3.1-3.3 (m, 2H), 3.6-3.8 (m, 2H), 4.3 (q, 2H), 4.6 (s, 2H), 6.9-7.3 (m, 7H), 7.5 (d, 1H).

5 Method 117

1,1-Dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a suspension of 1,1-dioxo-3,3-dipropyl-5-phenyl-7-bromo-8-methoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (prepared according to WO 96/16051 using identical synthetic steps except that the starting material was chosen to give the dipropyl compound instead of the butyl/ethyl compound; 0.756 g, 1.62 mmol) in DMF (40 ml) was added NaSMe (0.605 g, 8.20 mmol, 95 %), and the mixture was stirred over night at 120°C. The solvent was removed under reduced pressure and the residue was partitioned between EtOAc and 0.5 M HCl. The aqueous layer was extracted twice more with EtOAc and the combined organic extracts were dried (MgSO_4) and concentrated. The title compound was obtained in 0.665 g (98 %). NMR (500 MHz, DMSO- d_6) 0.60-0.80 (m, 6H), 1.05-1.50 (m, 8H), 2.15 (s, 3H), 3.20 (s, 2H), 3.65 (brs, 2H), 6.65 (s, 1H), 6.75-6.95 (m, 3H), 7.10-7.25 (m, 2H), 7.30 (s, 1H), 10.5 (s, 1H).

Method 118

20 1,1-Dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-carboxymethoxy-2,3,4,5-tetrahydro-1,5-benzothiazepine

To a suspension of 1,1-dioxo-3,3-dipropyl-5-phenyl-7-methylthio-8-hydroxy-2,3,4,5-tetrahydro-1,5-benzothiazepine (Method 117; 0.665 g, 1.58 mmol) in MeCN (10 ml) was added ethyl bromoacetate (0.262 ml, 2.35 mmol), tetrabutylammonium bromide (0.051 g, 0.158 mmol) and sodium carbonate (0.870 g, 8.21 mmol). The mixture was stirred over night at 80°C. The solvent was removed under reduced pressure and the residue was partitioned between EtOAc and 0.5 M HCl. The organic layer was washed with brine, dried (MgSO_4) and concentrated. The residue was filtered through a short silica column (DCM:EtOAc-9:1), concentrated and dissolved in EtOH (10 ml). A solution of NaOH (0.25 g, 6.25 mmol) in water (1 ml) was added and the solution was stirred over night at room temperature. The solvent was removed under reduced pressure and the residue was partitioned between EtOAc and 0.5 M HCl. The aqueous layer was extracted twice more with EtOAc and the combined organic extracts were washed with brine and concentrated. The crude product was purified by

preparative HPLC using a MeCN/ammonium acetate buffer to give the title compound in 0.441 g (58 %) as a white solid. NMR (DMSO-d_6) 0.55-0.75 (m, 6H), 1.05-1.50 (m, 8H), 2.15 (s, 3H), 3.20 (s, 2H), 3.65 (brs, 2H), 4.50 (s, 2H), 6.65 (s, 1H), 6.80-7.00 (m, 3H), 7.15 (s, 1H), 7.15-7.25 (m, 2H).

5

Example 121

The following illustrate representative pharmaceutical dosage forms containing the compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof (hereafter compound X), for therapeutic or prophylactic use in humans:-

(a): Tablet I	mg/tablet
Compound X	100
Lactose Ph.Eur	182.75
Croscarmellose sodium	12.0
Maize starch paste (5% w/v paste)	2.25
Magnesium stearate	3.0

10

(b): Tablet II	mg/tablet
Compound X	50
Lactose Ph.Eur	223.75
Croscarmellose sodium	6.0
Maize starch	15.0
Polyvinylpyrrolidone (5% w/v paste)	2.25
Magnesium stearate	3.0

(c): Tablet III	mg/tablet
Compound X	1.0
Lactose Ph.Eur	93.25
Croscarmellose sodium	4.0
Maize starch paste (5% w/v paste)	0.75
Magnesium stearate	1.0

(d): Capsule	mg/capsule
Compound X	10
Lactose Ph.Eur	488.5
Magnesium stearate	1.5

(e): Injection I	(50 mg/ml)
Compound X	5.0% w/v
1M Sodium hydroxide solution	15.0% v/v
0.1M Hydrochloric acid	(to adjust pH to 7.6)
Polyethylene glycol 400	4.5% w/v
Water for injection	to 100%

(f): Injection II	10 mg/ml
Compound X	1.0% w/v
Sodium phosphate BP	3.6% w/v
0.1M Sodium hydroxide solution	15.0% v/v
Water for injection	to 100%

(g): Injection III	(1 mg/ml, buffered to pH 6)
Compound X	0.1% w/v
Sodium phosphate BP	2.26% w/v
Citric acid	0.38% w/v
Polyethylene glycol 400	3.5% w/v
Water for injection	to 100%

5

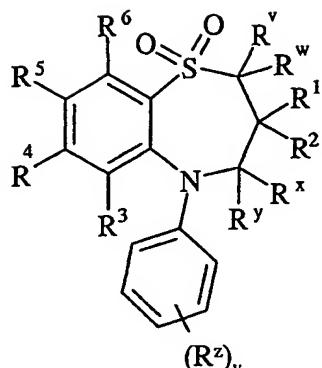
Note

The above formulations may be obtained by conventional procedures well known in the pharmaceutical art. The tablets (a)-(c) may be enteric coated by conventional means, for example to provide a coating of cellulose acetate phthalate.

Claims

What we claim is:

1. A compound of formula (I):



5

(I)

wherein:

R^v and **R^w** are independently selected from hydrogen or C₁₋₆alkyl;

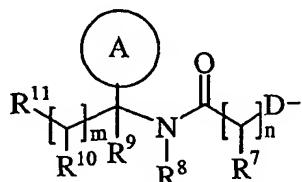
R¹ and **R²** are independently selected from C₁₋₆alkyl;

10 **R^x** and **R^y** are independently selected from hydrogen or C₁₋₆alkyl, or one of **R^x** and **R^y** is hydrogen or C₁₋₆alkyl and the other is hydroxy or C₁₋₆alkoxy;

R^z is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkanoyl, C₁₋₆alkanoyloxy, N-(C₁₋₆alkyl)amino, N,N-(C₁₋₆alkyl)₂amino, C₁₋₆alkanoylamino, N-(C₁₋₆alkyl)carbamoyl, 15 N,N-(C₁₋₆alkyl)₂carbamoyl, C₁₋₆alkylS(O)_a wherein a is 0 to 2, C₁₋₆alkoxycarbonyl, C₁₋₆alkoxycarbonylamino, ureido, N'-(C₁₋₆alkyl)ureido, N-(C₁₋₆alkyl)ureido, N',N'-(C₁₋₆alkyl)₂ureido, N'-(C₁₋₆alkyl)-N-(C₁₋₆alkyl)ureido, N,N'-(C₁₋₆alkyl)₂-N-(C₁₋₆alkyl)ureido, N-(C₁₋₆alkyl)sulphamoyl and N,N-(C₁₋₆alkyl)₂sulphamoyl;

20 **v** is 0-5;

 one of **R⁴** and **R⁵** is a group of formula (IA):



(IA)

R^3 and R^6 and the other of R^4 and R^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}$ alkyl)amino, $N,N-(C_{1-4}$ alkyl)₂amino, C_{1-4} alkanoylamino, $N-(C_{1-4}$ alkyl)carbamoyl,

5 $N,N-(C_{1-4}$ alkyl)₂carbamoyl, C_{1-4} alkylS(O)_a wherein a is 0 to 2, C_{1-4} alkoxycarbonyl, $N-(C_{1-4}$ alkyl)sulphamoyl and $N,N-(C_{1-4}$ alkyl)₂sulphamoyl; wherein R^3 and R^6 and the other of R^4 and R^5 may be optionally substituted on carbon by one or more R^{16} ;

D is -O-, -N(R^a)-, -S(O)_b- or -CH(R^a)-; wherein R^a is hydrogen or C_{1-6} alkyl and b is 0-2;

10 Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R^{17} ;

R^7 is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^7 is optionally substituted by one or more substituents selected from R^{18} ;

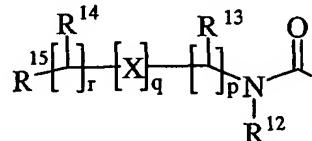
R^8 is hydrogen or C_{1-4} alkyl;

15 R^9 is hydrogen or C_{1-4} alkyl;

R^{10} is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^{10} is optionally substituted by one or more substituents selected from R^{19} ;

R^{11} is carboxy, sulpho, sulphino, phosphono, tetrazolyl, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently

20 selected from C_{1-6} alkyl; or R^{11} is a group of formula (IB):



(IB)

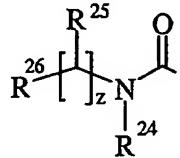
wherein:

X is $-N(R^q)-$, $-N(R^q)C(O)-$, $-O-$, and $-S(O)_a-$; wherein a is 0-2 and R^q is hydrogen or C_{1-4} alkyl;

R^{12} is hydrogen or C_{1-4} alkyl;

R^{13} and R^{14} are independently selected from hydrogen, C_{1-4} alkyl, carbocyclyl, heterocyclyl or R^{23} ; wherein said C_{1-4} alkyl, carbocyclyl or heterocyclyl may be independently optionally substituted by one or more substituents selected from R^{20} ;

R^{15} is carboxy, sulpho, sulphino, phosphono, tetrazolyl, $-P(O)(OR^e)(OR^f)$, $-P(O)(OH)(OR^e)$, $-P(O)(OH)(R^g)$ or $-P(O)(OR^e)(R^f)$ wherein R^e and R^f are independently selected from C_{1-6} alkyl; or R^{15} is a group of formula (IC):



5

(IC)

wherein:

R^{24} is selected from hydrogen or C_{1-4} alkyl;

R^{25} is selected from hydrogen, C_{1-4} alkyl, carbocyclyl, heterocyclyl or R^{27} ; wherein said C_{1-4} alkyl, carbocyclyl or heterocyclyl may be independently optionally substituted by one 10 or more substituents selected from R^{28} ;

R^{26} is selected from carboxy, sulpho, sulphino, phosphono, tetrazolyl, $-P(O)(OR^g)(OR^h)$, $-P(O)(OH)(OR^g)$, $-P(O)(OH)(R^g)$ or $-P(O)(OR^g)(R^h)$ wherein R^g and R^h are independently selected from C_{1-6} alkyl;

p is 1-3; wherein the values of R^{13} may be the same or different;

15 q is 0-1;

r is 0-3; wherein the values of R^{14} may be the same or different;

m is 0-2; wherein the values of R^{10} may be the same or different;

n is 1-3; wherein the values of R^7 may be the same or different;

z is 0-3; wherein the values of R^{25} may be the same or different;

20 R^{16} , R^{17} and R^{18} are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}$ alkyl)amino, $N,N-(C_{1-4}$ alkyl)₂amino, C_{1-4} alkanoylamino, $N-(C_{1-4}$ alkyl)carbamoyl, $N,N-(C_{1-4}$ alkyl)₂carbamoyl, C_{1-4} alkylS(O)_a wherein a is 0 to 2, C_{1-4} alkoxycarbonyl, $N-(C_{1-4}$ alkyl)sulphamoyl and

25 $N,N-(C_{1-4}$ alkyl)₂sulphamoyl; wherein R^{16} , R^{17} and R^{18} may be independently optionally substituted on carbon by one or more R^{21} ;

30 R^{19} , R^{20} , R^{23} , R^{27} and R^{28} are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}$ alkyl)amino, $N,N-(C_{1-4}$ alkyl)₂amino, C_{1-4} alkanoylamino, $N-(C_{1-4}$ alkyl)carbamoyl, $N,N-(C_{1-4}$ alkyl)₂carbamoyl, C_{1-4} alkylS(O)_a wherein a is 0 to 2, C_{1-4} alkoxycarbonyl, $N-(C_{1-4}$ alkyl)sulphamoyl,

N,N-(C₁₋₄alkyl)₂sulphamoyl, carbocyclyl, heterocyclyl, sulpho, sulphino, amidino, phosphono, -P(O)(OR^a)(OR^b), -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or -P(O)(OR^a)(R^b), wherein R^a and R^b are independently selected from C₁₋₆alkyl; wherein R¹⁹, R²⁰, R²³, R²⁷ and R²⁸ may be independently optionally substituted on carbon by one or more R²²;

5 **R²¹ and R²²** are independently selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl, formamido, acetylamino, acetoxy, methylamino, dimethylamino, *N*-methylcarbamoyl, *N,N*-dimethylcarbamoyl, methylthio, methylsulphanyl, mesyl, *N*-methylsulphamoyl and
10 *N,N*-dimethylsulphamoyl;
or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

2. A compound of formula (I) according to claim 1 wherein R^v and R^w are both hydrogen or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

15 3. A compound of formula (I) according to either of claims 1 or 2 wherein R¹ and R² are independently selected from ethyl, propyl or butyl or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

20 4. A compound of formula (I) according to any one of claims 1 to 3 wherein R^x and R^y are both hydrogen or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

5. A compound of formula (I) according to any one of claims 1 to 4 wherein R^z is
25 selected from halo, amino, C₁₋₆alkyl, C₁₋₆alkoxycarbonylamino or *N'*-(C₁₋₆alkyl)ureido or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

30 6. A compound of formula (I) according to any one of claims 1 to 5 wherein v is 0 or 1 or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

7. A compound of formula (I) according to any one of claims 1 to 6 wherein R³ is hydrogen or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

8. A compound of formula (I) according to any one of claims 1 to 7 wherein the R⁴ or R⁵ that is not the group of formula (IA) is selected from hydrogen, halo, C₁₋₄alkoxy or C₁₋₄alkylS(O)_a wherein a is 0 to 2; wherein that R⁴ or R⁵ may be optionally substituted on 5 carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy, carboxy and N,N-(C₁₋₄alkyl)₂amino or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

9. A compound of formula (I) according to any one of claims 1 to 8 wherein R⁵ is a 10 group of formula (IA) (as depicted in claim 1) and R⁴ is methylthio or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

10. A compound of formula (I) according to any one of claims 1 to 9 wherein R⁶ is 15 hydrogen or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

11. A compound of formula (I) according to any one of claims 1 to 10 wherein in the group of formula (IA):

D is -O- or -S-;

20 Ring A is phenyl, thienyl or indolyl; wherein Ring A is optionally substituted by one or more substituents selected from halo, hydroxy, methoxy or trifluoromethyl;

R⁷ is hydrogen, methyl or phenyl;

R⁸ is hydrogen or methyl;

R⁹ is hydrogen or methyl;

25 R¹⁰ is hydrogen;

m is 0-2 wherein the values of R¹⁰ may be the same or different; and

R¹¹ is carboxy, -P(O)(OH)(OEt) or a group of formula (IB) (as depicted in claim 1); or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

30 12. A compound of formula (I) according to any one of claims 1 to 11 wherein in the group of formula (IB):

R¹² is hydrogen or methyl;

R^{13} is hydrogen, methyl, ethyl, butyl or phenyl or R^{23} ; wherein R^{13} is optionally substituted by one or more substituents selected from R^{20} ; R^{20} is hydroxy, methylthio, methoxy, amino, imidazolyl or mercapto; wherein R^{20} may be independently optionally substituted on carbon by one or more hydroxy; R^{23} is carboxy;

5 X is $-NH-$ or $-NHC(O)-$;

R^{14} is selected from hydrogen, methyl or phenyl; wherein said methyl or phenyl may be optionally substituted by one or more substituents selected from hydroxy;

10 R^{15} is carboxy, sulpho, phosphono, $-P(O)(OR^e)(OR^f)$, $-P(O)(OH)(OR^e)$, $-P(O)(OH)(R^e)$ or $-P(O)(OR^e)(R^f)$ wherein R^e and R^f are independently selected from methyl or ethyl or R^{15} is a group of formula (IC) (as depicted in claim 1);

p is 1-3 wherein the values of R^{13} may be the same or different;

q is 0-1; and

r is 0-3 wherein the values of R^{14} may be the same or different;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

15

13. A compound of formula (I) according to any one of claims 1 to 12 wherein in the group of formula (IC):

R^{24} is hydrogen;

R^{25} is hydrogen;

20 R^{26} is carboxy; and

z is 1;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

14. A compound of formula (I) (as depicted in claim 1) wherein:

25 R^v and R^w are both hydrogen;

R^1 and R^2 are independently selected from C_{1-4} alkyl;

R^x and R^y are both hydrogen;

R^z is selected from halo, amino, C_{1-6} alkyl, C_{1-6} alkoxycarbonylamino or

$N'-(C_{1-6}$ alkyl)ureido;

30 v is 0 or 1;

R^3 and R^6 are hydrogen;

 one of R^4 and R^5 is a group of formula (IA) (as depicted in claim 1) and the other is selected from hydrogen, halo, C_{1-4} alkoxy or C_{1-4} alkylS(O)_a wherein a is 0 to 2; wherein that

R⁴ or R⁵ may be optionally substituted on carbon by one or more R¹⁶; wherein R¹⁶ is independently selected from hydroxy, carboxy and N,N-(C₁₋₄alkyl)₂amino;

D is -O- or -S-;

R⁷ is hydrogen, methyl or phenyl;

5 R⁸ is hydrogen or methyl;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R¹⁷; wherein R¹⁷ is selected from halo, hydroxy, C₁₋₄alkyl or C₁₋₄alkoxy; wherein R¹⁷ may be optionally substituted on carbon by one or more R²¹; wherein R²¹ is selected from halo;

10 R⁹ is hydrogen or methyl;

R¹⁰ is hydrogen;

R¹¹ is carboxy, -P(O)(OH)(OR^c) wherein R^c is selected from C₁₋₄alkyl or a group of formula (IB) (as depicted in claim 1);

R¹² is hydrogen or methyl;

15 X is -NH- or -NHC(O)-;

R¹³ is hydrogen, C₁₋₄alkyl, carbocyclyl or R²³; wherein R¹³ is optionally substituted by one or more substituents selected from R²⁰; wherein R²⁰ is hydroxy, C₁₋₄alkylS(O)_a wherein a is 0, C₁₋₄alkoxy, amino, carbocyclyl, heterocyclyl or mercapto; wherein R²⁰ may be independently optionally substituted on carbon by one or more R²²; R²² is selected from hydroxy; and R²³ is carboxy;

R¹⁴ is selected from hydrogen, C₁₋₄alkyl or carbocyclyl; wherein said C₁₋₄alkyl or carbocyclyl may be optionally substituted by one or more substituents selected from R²⁰; and R²⁰ is hydroxy;

R¹⁵ is carboxy, sulpho, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e),

25 -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₄alkyl or R¹⁵ is a group of formula (IC) (as depicted in claim 1);

R²⁴ is hydrogen;

R²⁵ is hydrogen;

R²⁶ is carboxy;

30 p is 1-3; wherein the values of R¹³ may be the same or different;

q is 0-1;

r is 0-3; wherein the values of R¹⁴ may be the same or different;

m is 0-2; wherein the values of R¹⁰ may be the same or different;

n is 1-2; wherein the values of R⁷ may be the same or different;
z is 0-1; wherein the values of R²⁵ may be the same or different;
or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

5 15. A compound of formula (I) (as depicted in claim 1) selected from:

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(carboxymethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(carboxymethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

10 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

15 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

20 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-carboxyethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(5-carboxypentyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

25 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{ α -[N'-(2-sulphoethyl)carbamoyl]-2-fluorobenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

30 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{*N*-[(R)- α -(*N'*-{(R)-1-[*N''*-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]-2-hydroxyethyl}carbamoyl)benzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{ α -[*N'*-(carboxymethyl)carbamoyl]

5 benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{ α -[*N'*-((ethoxy)(methyl)phosphoryl)methyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-{*N*-[(R)- α -(*N'*-{2-[(hydroxy)(methyl)phosphoryl]ethyl}carbamoyl)benzyl]carbamoylmethoxy}-2,3,4,5-

10 tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-{(2-methylthio-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{*N*-[(R)- α -(*N'*-{2-[(methyl)(ethyl)phosphoryl]ethyl}carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-

15 benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{*N*-[(R)- α -(*N'*-{2-[(methyl)(hydroxy)phosphoryl]ethyl}carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[(R)-*N'*-(2-methylsulphinyl-1-

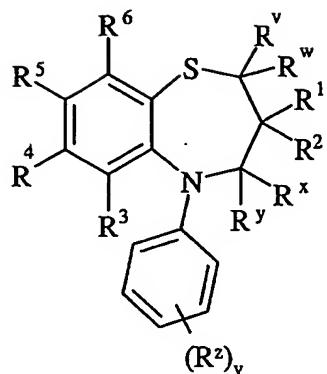
20 carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine; and

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-[*N*-{(R)- α -[(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine; or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

25

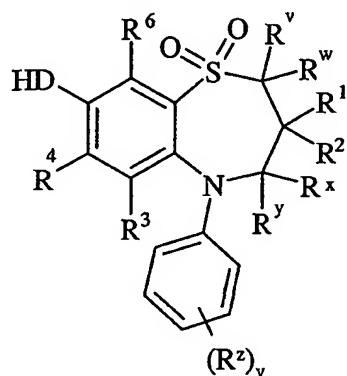
16. A process for preparing a compound of the formula (I) as claimed in any one of claims 1 to 15 which comprises of:-

Process 1): oxidising a benzothiazepine of formula (II):



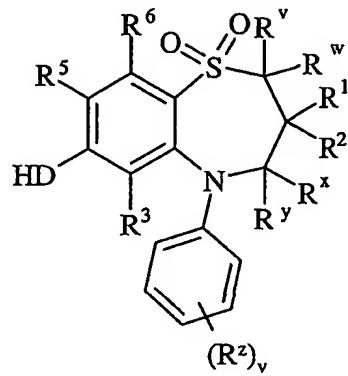
(III);

Process 2): for compounds of formula (I) wherein D is -O-, -NR^a or -S-; reacting a compound of formula (IIIa) or (IIIb):



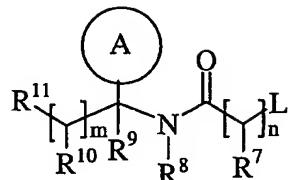
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(IIIa)



(IIIb)

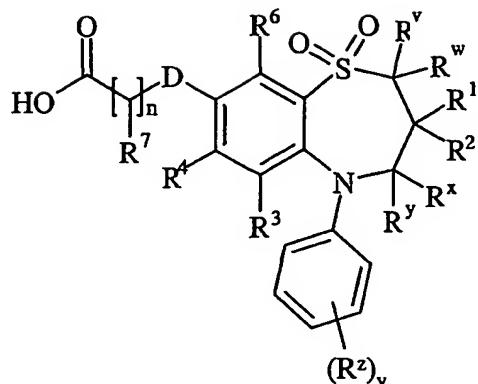
with a compound of formula (IV):



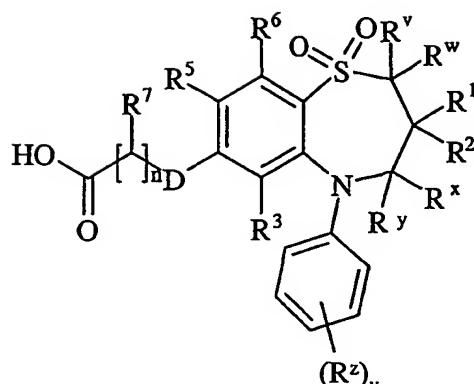
(IV)

10 wherein L is a displaceable group;

Process 3): reacting an acid of formula (Va) or (Vb):

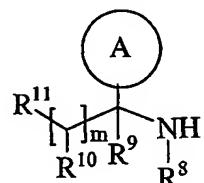


(Va)



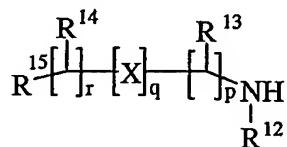
(Vb)

or an activated derivative thereof; with an amine of formula (VI):



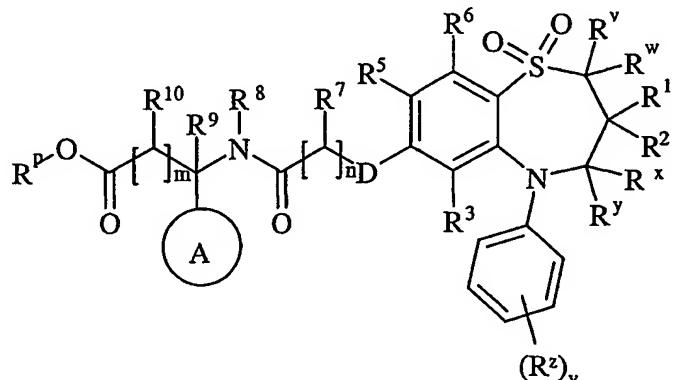
(VI);

Process 4): for compounds of formula (I) wherein R¹¹ is a group of formula (IB); reacting a compound of formula (I) wherein R¹¹ is carboxy with an amine of formula (VII):



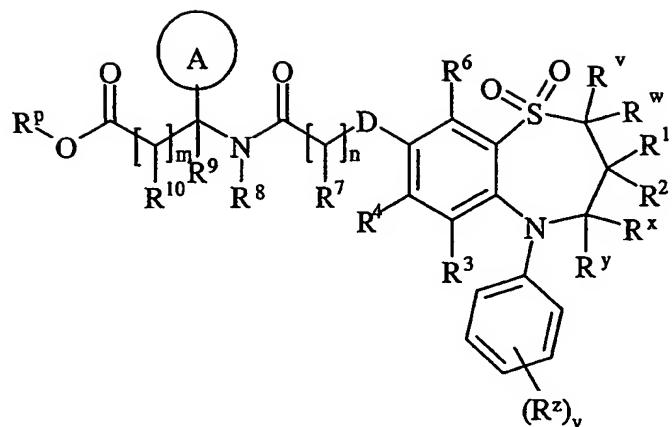
(VII)

10 Process 5): for compounds of formula (I) wherein R¹¹ is carboxy; deprotecting a compound of formula (VIIIa):



(VIIIa)

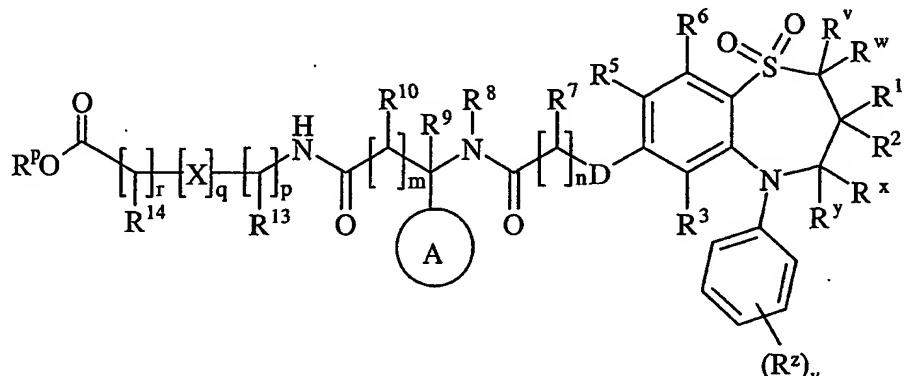
or (VIIIb):



(VIIIb)

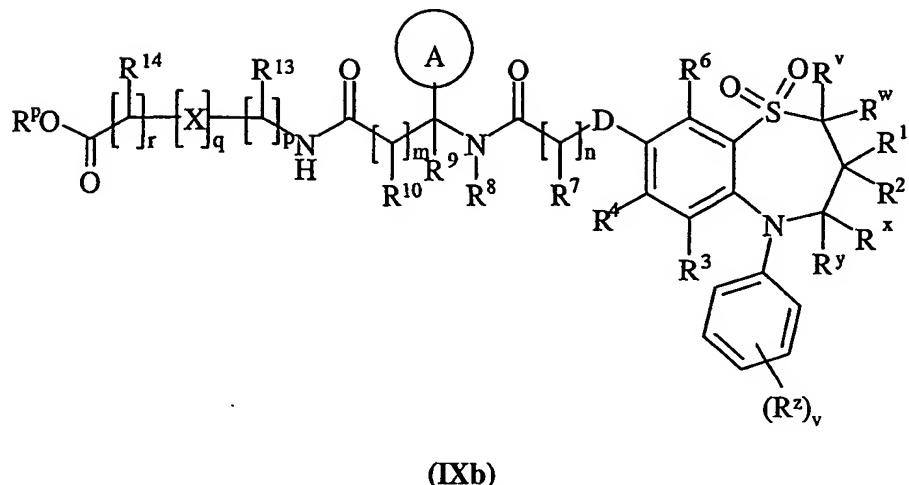
wherein R^p is C_{1-4} alkyl;

5 *Process 6*): for compounds of formula (I) wherein R^{11} is a group of formula (IB) and R^{15} is carboxy; deprotecting a compound of formula (IXa):



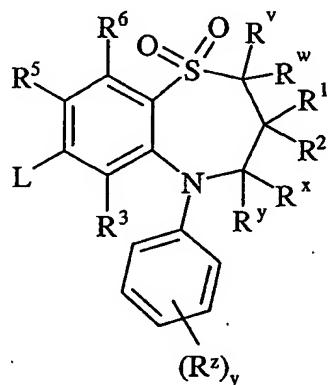
(IXa)

10 or (IXb):

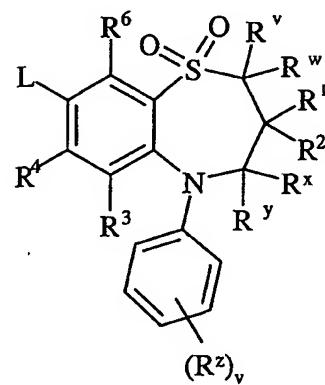


wherein R^p is C_{1-4} alkyl;

Process 7) for compounds of formula (I) wherein one of R^4 and R^5 are independently selected from C_{1-4} alkylthio optionally substituted on carbon by one or more R^{16} ; reacting a compound of formula (Xa) or (Xb):



(Xa)



(Xb)

10 wherein L is a displaceable group; with a thiol of formula (XI):

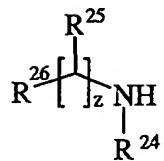


(XI)

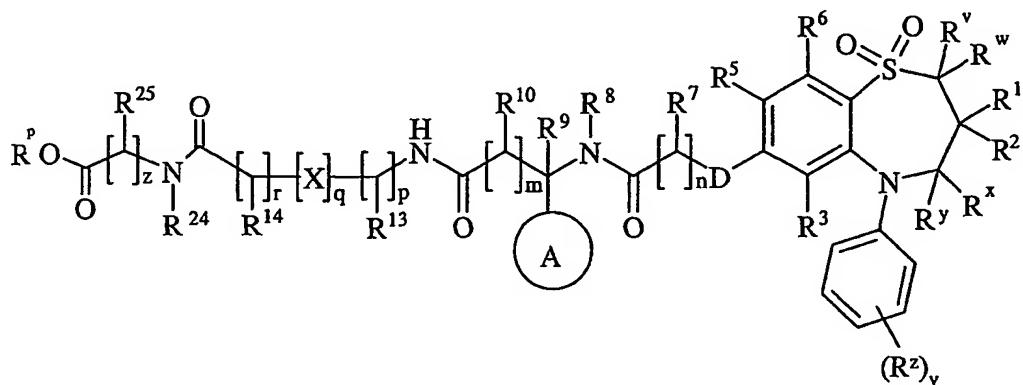
wherein R^y is C_{1-4} alkylthio optionally substituted on carbon by one or more R^{16} ;

Process 8) for compounds of formula (I) wherein R^{15} is a group of formula (IC) reacting a compound of formula (IXa) or (IXb) wherein R^p is hydrogen with a compound of formula (XII):

- 154 -

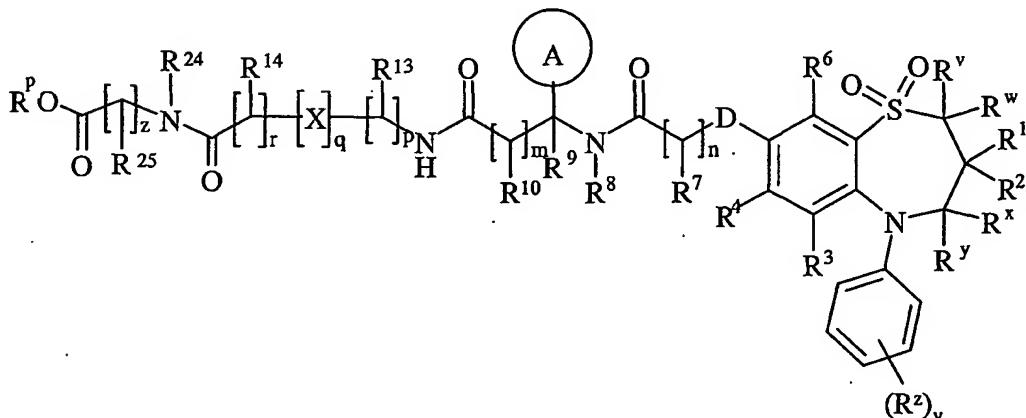


Process 9): for compounds of formula (I) wherein R¹¹ is a group of formula (IB) and R¹⁵ is a group of formula (IC) and R²⁶ is carboxy; deprotecting a compound of formula (XIIIa):



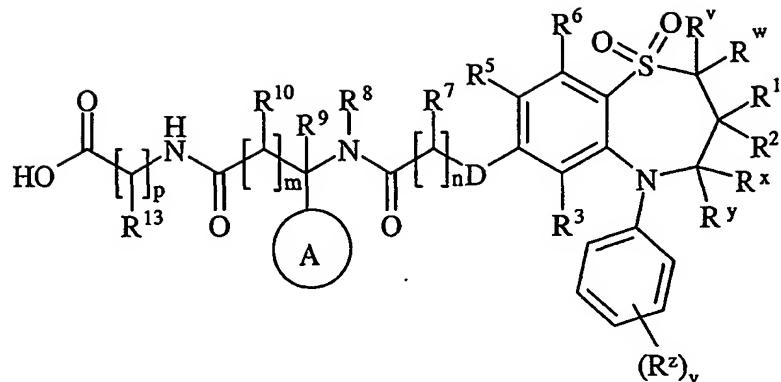
5

or (XIIIb):

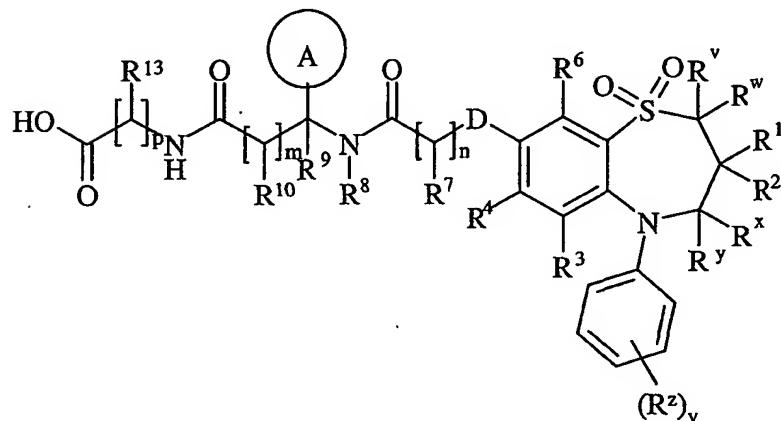


10 and R^p is C₁₋₄alkyl;

Process 10): for compounds of formula (I) wherein X is -N(R⁹)C(O)-; reacting a compound of formula (XIVa):

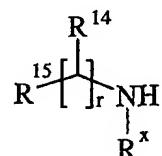


or (XIVb):



5

with a compound of formula (XV):



and thereafter if necessary or desirable:

- 10 i) converting a compound of the formula (I) into another compound of the formula (I);
- ii) removing any protecting groups;
- iii) forming a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug.

17. A compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15 for use as a medicament.

18. A compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15 for use in a method of prophylactic or therapeutic treatment of a warm-blooded animal, such as man.

5

19. The use of a compound of the formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15 in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

10

20. A method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15.

15

21. A pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15, in association with a pharmaceutically-acceptable diluent or carrier.

20

22. A pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in association with a pharmaceutically acceptable diluent or carrier.

25

23. A pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15, and a bile acid binder, in association with a pharmaceutically acceptable diluent or carrier.

30

24. A pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as

claimed in any one of claims 1 to 15, and an HMG Co-A reductase inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a bile acid binder in association with a pharmaceutically acceptable diluent or carrier.

5. 25. A composition according to claim 22 or claim 24 wherein the HMG Co-A reductase inhibitor is atorvastatin, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

26. A composition according to claim 22 or claim 24 wherein the HMG Co-A reductase
10 inhibitor is rosuvastatin, or a pharmaceutically acceptable salt thereof.

27. A pharmaceutical composition which comprises a compound of formula (I), or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, as claimed in any one of claims 1 to 15 and a PPAR alpha and/or gamma agonist, or a
15 pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable diluent or carrier.

28. A composition according to claim 27 wherein the PPAR alpha and/or gamma agonist is (S)-2-ethoxy-3-[4-(2-{4-methanesulphonyloxyphenyl}ethoxy)phenyl]propanoic acid or a
20 pharmaceutically acceptable salt thereof.

29. A compound of formula (VIIIa), (VIIIb), (IXa), (IXb), (XIIIa) or (XIIIb) (as depicted in claim 16) or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

INTERNATIONAL SEARCH REPORT

PCT/GB 01/05554

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07D281/10 C07D417/12 C07F9/6536 C07K5/065 C07K5/087 A61K31/554 A61K31/675 A61K38/05 A61K38/06 A61P3/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D C07F C07K A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, BEILSTEIN Data, WPI Data, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 99 35135 A (GLAXO GROUP LIMITED) 15 July 1999 (1999-07-15) cited in the application the whole document ---	1
A	WO 96 16051 A (THE WELLCOME FOUNDATION LIMITED) 30 May 1996 (1996-05-30) cited in the application the whole document ---	1
P, X	WO 01 66533 A (ASTRAZENECA AB ET AL) 13 September 2001 (2001-09-13) the whole document ---	1-29

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- *&* document member of the same patent family

Date of the actual completion of the international search

28 February 2002

Date of mailing of the international search report

08/03/2002

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Authorized officer

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FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1-29 (all partly)

The scope of claims 1-29, in as far as the expression "prodrugs" is concerned, is so unclear (Article 6 PCT) that a meaningful international search is impossible with regard to this expression.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

PCT/GB 01/05554

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